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04/22/98

SHERIDAN NEIMARK  
ROGER L BROWDY

ANNE M KORNBAU  
NORMAN J LATKER  
NICK BROMER\*  
(PA BAR ONLY)

OF COUNSEL  
IVER P COOPER

# BROWDY AND NEIMARK, PLLC

ATTORNEYS AT LAW  
PATENT AND TRADEMARK CAUSES

SUITE 300  
419 SEVENTH STREET, N W  
WASHINGTON, D. C 20004-2299

TELEPHONE (202)-628-5197

April 22, 1998

TELECOPIER FACSIMILE  
(202) 737-3528  
(202) 393-1012

E-MAIL  
BrwdyNmrk@digizen.net

SENIOR COUNSEL  
ALVIN BROWDY

PATENT AGENT  
ALLEN C YUN, PHD

Hon. Assistant Commissioner for Patents  
Box Patent Appln  
Washington, D.C. 20231

RE: New Patent Application in U.S.  
Applicant(s): Toshio ARIYASU et al.  
Title: HEDGEHOG PROTEIN  
Atty's Docket: ARIYASU=1

Sir:

Attached herewith is the above-identified application for Letters Patent including:

- ☒ Specification (67 pages), claims (3 pages) and abstract (1 page)
- ☒ 3 Sheets Drawings (Figures 1-3)
  - ☒ Formal ☐ Informal
- ☒ Declaration and Power of Attorney (2 pages)
  - ☒ Newly executed ☐ Copy from prior application no.
- ☐ Preliminary Amendment
- ☐ Supplemental Preliminary Amendment
- ☐ Information Disclosure Statement with ( ) references
- ☐ A verified statement to establish small entity status under 37 CFR §1.9 and 37 CFR §1.27 ( page(s))
- ☒ A check in the amount of \$ 834.00 (check no. 18559) to cover:
  - ☒ The filing fee calculated as follows:

CLAIMS AS FILED				
FOR	NUMBER FILED	NUMBER EXTRA	RATE	BASIC FEE \$ 790.00
TOTAL CLAIMS	22 - 20=	2	x 22	\$ 44.00
INDEPENDENT CLAIMS	1 - 3=	0	x 82	0
<input type="checkbox"/> Multiple Dependent Claim Presented			x270	
<input type="checkbox"/> Reduction of 1/2 for small entity				- \$
			TOTAL FILING FEE	\$ 834.00

- ☐ Any additional fee required by the filing of an enclosed preliminary or supplemental preliminary amendment has been calculated as shown below:

	CLAIMS REMAINING AFTER AMENDMENT	HIGHEST NO. PREVIOUSLY PAID FOR	PRESENT EXTRA	RATE	CALCULATION
TOTAL	*	MINUS **	=	x \$ 22.00	\$
INDEP	*	MINUS ***	=	x \$ 82.00	\$
[ ] Multiple Dependent Claim Presented				x \$270.00	\$
Total of Above Calculations =					\$
Reduction by 1/2 for filing by small entity					- \$
Total Additional Fee =					\$

[ ] \_\_\_\_\_.

[X] Return Receipt Postcard (in duplicate)

The following statements are applicable:

[X] The benefit under 35 USC §119 is claimed of the filing date of: Japanese Application No. 121578/1997 in Japan on April 25, 1997; and Japanese Application No. not yet received in Japan on April 14, 1998. A certified copy of said priority documents

[ ] is attached

[ ] was filed in progenitor case \_\_\_\_\_ on \_\_\_\_\_.

Application No. \_\_\_\_\_ in \_\_\_\_\_ on \_\_\_\_\_. A certified copy of said priority document [ ] is attached [ ] was filed in progenitor case \_\_\_\_\_ on \_\_\_\_\_.

[ ] The present application is a [ ] Continuation [ ] Divisional

[ ] Continuation-in-part of prior application No. \_\_\_\_\_.

[ ] Incorporation By Reference. The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied herewith, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein.

[ ] A signed statement deleting inventor(s) named in the prior application is attached.

[ ] Amend the specification by inserting before the first line the sentence: --This is a \_\_ continuation \_\_ division of copending parent application Serial No. filed \_\_\_\_\_.--

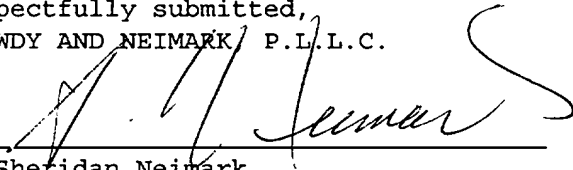
[ ] Certain documents were previously cited or submitted to the Patent and Trademark Office in the following prior application \_\_\_\_\_, which is relied upon under 35 USC §120. Applicants identify these documents by attaching hereto a form PTO-1449 listing these documents, and request that they be considered and made of record in accordance with 37 CFR §1.98(d). Per Section 1.98(d), copies of these documents need not be filed in this application.

[ ] A verified statement claiming small entity status is enclosed in progenitor application no. \_\_\_\_\_, filed \_\_\_\_\_. Status is still proper and desired.

[ ] The undersigned attorney of record hereby revokes the powers of attorney of:

- [ ] The undersigned attorney of record hereby appoints associate power of attorney, to prosecute this application and to transact all business in the Patent and Trademark Office in connection therewith to:
- [X] The Commissioner is hereby authorized to charge payment of the following additional fees associated with this communication or credit any overpayments to Deposit Account No. 02-4035:
- [X] Any additional filing fees required under 37 CFR §1.16.
- [X] Any patent application processing fees under 37 CFR §1.17.
- [X] The Commissioner is hereby authorized to charge payment of the following fees, based on any paper filed during the pendency of this application or any CPA thereof, to effect any amendment, petition, or other action requested in said paper or credit any overpayments to Deposit Account No. 02-4035:
- [X] Any patent application processing fees under 37 CFR §1.17.
- [ ] The issue fee set in 37 CFR §1.18 at or before mailing the Notice of Allowance, pursuant to 37 CFR §1.311(b).
- [X] Any filing fees under 37 CFR §1.16 for presentation of extra claims.
- [X] If a paper is untimely filed in this or any CPA thereof by Applicant(s), the Commissioner is hereby petitioned under 37 CFR §1.136(a) for the minimum extension of time required to make said paper timely. In the event a petition for extension of time is made under the provisions of this paragraph, the Commissioner is hereby requested to charge any fee required under 37 CFR §1.17 to Deposit Account 02-4035.
- [X] The Commissioner is hereby authorized to credit any overpayment of fees accompanying this paper to Deposit Account No. 02-4035.

Respectfully submitted,  
BROWDY AND NEIMARK/ P.L.L.C.

By:   
Sheridan Neimark  
Registration No. 20,520

SN:bcs

f:\user3\98apr\arl.ctr

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:	)	Art Unit:
	)	
Toshio ARIYASU et al.	)	Washington, D.C.
	)	
U.S.S.N.:	)	April 22, 1998
	)	
Filing Date: 22 April 1998	)	
	)	
For: HEDGEHOG PROTEIN	)	Docket No.: ARIYASU=1

PRELIMINARY AMENDMENT

Honorable Commissioner of Patents and Trademarks  
Washington, D.C. 20231

Sir:

Contemporaneous with the filing of this case and prior to calculation of the filing fee, kindly amend as follows:

IN THE CLAIMS

Claim 18, line 3, delete "of claim 1".

If, inadvertently, a proper multiple dependent claim has not been amended to reduce it to single dependency, please amend it to be dependent solely on the first-mentioned claim, or, if that is not possible, please cancel the claim and notify the undersigned.

REMARKS

The above amendment to the claims is being made in order to eliminate multiple dependency and for the purpose of reducing the filing fee. Please enter this amendment prior to calculation of the filing fee in this case.

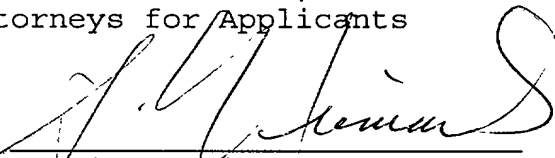
In re of ARIYASU=1

Favorable consideration and allowance are earnestly  
solicited.

Respectfully submitted,

BROWDY AND NEIMARK, P.L.L.C.  
Attorneys for Applicants

By

  
Sheridan Neimark  
Registration No. 20,520

SN:bcs  
Telephone No.: (202) 628-5197  
Facsimile No.: (202) 737-3528

f:\user3\98apr\anl.pa

## HEDGEHOG PROTEIN

### Background of the Invention

#### 1. Field of the Invention

This invention relates to a novel hedgehog protein, more particularly, a Desert hedgehog protein of human origin.

#### 2. Description of the Prior Art

The hedgehog gene was originally identified by genetic techniques as a gene that plays an important role in normal morphogenesis during embryonic and larval development in the fruit fly *Drosophila melanogaster*, as described by C. Nüsslein-Volhard et al., in *Nature*, Vol.287, pp.795-801 (1980). J. J. Lee et al. sequenced the gene and deduced the amino acid sequence of the hedgehog protein as the expression product in *Cell*, Vol.71, pp.33-50 (1992). Some homologues of the gene were later isolated from vertebrates including mammals (hereinafter, the homologues of species other than the fruit fly may also be called "hedgehog"). At present, it is known that vertebrate hedgehog genes, unlike that of the fruit fly, form a multigene family and would play different roles in normal morphogenesis.

For example, as described by Y. Echelard et al., in *Cell*, Vol.75, pp.1417-1430 (1993), there have been identified three types of the genes of mouse origin, designated "Sonic hedgehog", "Indian hedgehog", and "Desert hedgehog", which have different nucleotide sequences and express in different manners in living bodies. While in human, there have been found only two types of the genes designated "Sonic hedgehog" and "Indian hedgehog", as described by V. Marigo et al., in *GENOMICS*, Vol.28,

pp.44-51 (1995), and their expression manners and functions of their expression products remain to be elucidated. Therefore from scientific and pharmaceutical viewpoints, in order to elucidate the process of exhibiting hereditary morphological abnormalities in humans and direct for their treatments and diagnoses, the establishment of a novel hedgehog gene and its expression product, i.e., a novel hedgehog protein are now in great expectation.

#### Summary of the Invention

In view of the foregoing, the first object of this invention is to provide a novel hedgehog protein of human origin.

The second object of this invention is to provide a DNA encoding the hedgehog protein.

The third object of this invention is to provide a monoclonal antibody recognizing the hedgehog protein.

The fourth object of this invention is to provide a process for producing the hedgehog protein.

The fifth object of this invention is to provide a method for detecting the hedgehog protein.

The present inventors energetically and extensively screened for human cell lines which express a novel hedgehog gene capable of attaining the above objects by using RT-PCR techniques, where RNAs obtained from various established human cell lines were used as templates, while as primers various oligonucleotides were chemically synthesized based on the nucleotide sequence of mouse Desert hedgehog gene registered in "GenBank®", a nucleic acid database by National Institute of

Health, USA, under the accession number "X76292". These screenings resulted in finding that some human cell lines including ARH-77 cell, ATCC CRL-1621, a cell line derived from plasma cell of a leukemia patient, expressed a specific gene in an elevated level checked with the above RT-PCRs. Further energetic studies confirmed that the human gene was a novel gene, which contained no nucleotide sequences that he said to know. Comparison with other genes revealed that the human gene has a relatively high homology to mouse Desert hedgehog gene. These findings led to the conclusion that the gene is a novel type of Desert hedgehog gene of human origin. A DNA obtained from the gene thus sequenced was introduced into *Escherichia coli* using an autonomously replicable vector, attaining satisfactory DNA expression and production of human Desert hedgehog protein.

Furthermore, the present inventors prepared known human Sonic hedgehog protein by using conventional recombinant DNA techniques and prepared monoclonal antibodies recognizing the protein. It was found that some of the monoclonal antibodies unexpectedly recognized well not only human Sonic hedgehog protein but also human Desert hedgehog protein. This invention was established based on these findings.

More particularly, the first object of this invention is attained by a Desert hedgehog protein of human origin.

The second object of this invention is attained by a DNA which encodes the hedgehog protein.

The third object of this invention is attained by a monoclonal antibody which recognizes the hedgehog protein.

The forth object of this invention is attained by a process for preparing a hedgehog protein which comprises the



steps of allowing to express a DNA encoding the hedgehog protein and collecting the generated hedgehog protein.

The fifth object of this invention is attained by a method for detecting the hedgehog protein using a monoclonal antibody which recognizes the hedgehog protein.

#### Brief Explanation of the Accompanying Drawings

FIG. 1 is the restriction map of the recombinant DNA of this invention "pHuDHH/pGEX-2T/#4-8".

FIG. 2 is a half tone image of gel electrophoresis given on a display, visualized by Western blotting as the detection method using the monoclonal antibody of this invention.

FIG. 3 shows the results of detecting the hedgehog protein by an enzyme-immunoassay as the detection method using the monoclonal antibody of this invention.

In FIG. 1, the symbol "HuDHH" indicates a DNA encoding the hedgehog protein of this invention; the symbol "Amp", an ampicillin-resistant gene; the symbol "pBR322ori", a replication origin exerting in *Escherichia coli*; the symbol "GST", a structural gene of glutathione S-transferase; and the symbol "Ptac", a Tac promotor.

#### Detailed Description of the Invention

This invention relates to a novel hedgehog protein, more particularly, a Desert hedgehog protein of human origin. The present hedgehog protein contains a part or the whole of the

amino acid sequence of SEQ ID NO:1, which may bear a homology, usually about 80%, to mouse Desert hedgehog protein at amino acid sequence level. Examples of the present hedgehog protein are a mature form of human Desert hedgehog protein that contains the amino acid sequence of SEQ ID NO:1 and a precursor form of human Desert hedgehog protein that has the amino acid sequence of SEQ ID NO:2 or 3, which contains the amino acid sequence of SEQ ID NO:1. The present hedgehog protein further includes other types of proteins with amino acid sequences as illustrated above where one or more amino acids are deleted or replaced with other ones, one or more amino acids are added, or saccharide chains are linked, so far as they contain the amino acid sequence as mentioned above. The present hedgehog protein shall not be restricted to those obtained from specific sources and by specific preparation methods, therefore it includes natural proteins obtained from cultures of established cell lines, recombinant proteins obtained by recombinant DNA techniques, and synthetic polypeptides obtained by way of peptide synthesis.

The DNA of this invention includes any DNAs which encode such hedgehog protein, regardless of their sources or origins. Thus the DNA of this invention includes those from natural sources as well as those artificially modified or chemically synthesized, as far as they encode the hedgehog protein of this invention. Generally in this field, in case of artificially expressing DNAs which encode proteins, one may replace one or more nucleotides in the DNAs with different nucleotides and/or link appropriate nucleotide sequences thereto with the purpose of improving their expression efficiency and/or the physiological and physicochemical properties of the protein.

Such modification are feasible in the DNA of this invention. More particularly, one can link, for example, to the 5'- and/or 3'-termini of the DNA as described above, recognition sites for appropriate restriction enzymes, initiation codons, termination codons, promoters and/or enhancers, as far as the final protein products do retain prescribed properties. Thus, the wording "DNA" as referred to in this invention shall mean, in addition to those which encode the above-mentioned proteins, those which are complementary thereto, and those where one or more nucleotides have been replaced with different nucleotides while conserving the encoding amino acid sequence.

Such a DNA can be obtained from natural by screening of human cells, for example, mammalian cells including epithelial cells, endothelial cells, interstitial cells, chondrocytes, monocytes, granulocytes, lymphocytes, neurocytes, and established cell lines from them of human origin, based on a hybridization with a DNA as a probe which encodes at least a part of the amino acid sequence of human Desert hedgehog protein of this invention, for example, the amino acid sequence of SEQ ID NO:1. Such screening can be achieved with conventional methods commonly used in this field such as PCR, RT-PCR, screening cDNA libraries, screening genomic libraries and/or modified methods thereof. Examples of preferred cells are established cell lines including ARH-77 cell, ATCC CRL-1621, K-562 cell, ATCC CCL-243, and KU-812 cell, an cell line reported by K. Kishi, in *Leukemia Research*, Vol.9, pp.381-390 (1985), and bone marrow cells. The DNA of this invention thus obtained usually contains a part or the whole of the nucleotide sequence of SEQ ID NO:4. For example, from ARH-77 cell, ATCC CRL-1621, a DNA encoding a mature form of human Desert

hedgehog protein that contains the nucleotide sequence of SEQ ID NO:4 or a DNA encoding a precursor form of human Desert hedgehog protein that has the nucleotide sequence of SEQ ID NO:5 or 6, which contains the nucleotide sequence of SEQ ID NO:4, can be obtained. The present DNA can also be obtained by conventional chemical synthesis. The DNA of this invention, once obtained in any manner, can be easily amplified to desired level by methods of PCR or those using autonomously replicable vectors.

The DNA of this invention includes those in the forms of recombinant DNAs where the DNA, encoding the present hedgehog protein, is inserted into autonomously replicable vectors. The recombinant DNAs can be relatively-easily obtained by using conventional recombinant DNA techniques, once the desired DNA is obtained. Examples of the vectors feasible in this invention are plasmid vectors including pGEX-2T, pGEX-4T-1, pKK223-3, pcDNAI/Amp, BCMGSNeo, pcDL-SR $\alpha$ , pKY4, pCDM8, pCEV4, and pME18S. The autonomously replicable vectors usually comprise nucleotide sequences suitable for the DNA expression in respective hosts, for example, promoters, enhancers, replication origins, terminators for transcription, splicing sequences, and/or sequences for selection markers. As the promotor, using a heat shock protein promotor or the interferon- $\alpha$  promotor disclosed in Japanese Patent Kokai No.163,368/95 by the same applicant makes it possible to regulate the present DNA expression in the transformants by external stimuli.

To insert the DNA of this invention, conventional methods commonly used in this field can be used. More particularly, a gene containing the DNA of this invention and an autonomously replicable vector are first cleaved with restriction

enzymes and/or ultrasonication, then the resulting DNA and vector fragments are ligated. Ligation of the DNA and vector fragments become much easier when the genes and vectors are digested with restriction enzymes specific to particular nucleotides, for example, *AccI*, *BamHI*, *BstXI*, *EcoRI*, *HindIII*, *NotI*, *PstI*, *SacI*, *Sall*, *SmaI*, *SpeI*, *XbaI* and *XhoI*. To ligate the DNA and vector fragments, they can be first annealed, if necessary, and then exposed to DNA ligase *in vivo* or *in vitro*. The recombinant DNAs thus obtained are unlimitedly replicable in hosts of microbe and animal origins.

The DNA of this invention further includes those in the forms where the DNA, encoding the above-mentioned hedgehog protein, is introduced into desired hosts. The DNA in such forms can be obtained without considerable difficulty by introducing the recombinant DNA of this invention to desired hosts. For the hosts, cells of microbe, animal or plant origin conventionally used in this field can be arbitrarily used. The use of the hosts of microbe origin has a merit of a higher productivity of the protein per culture. The hosts of animal origin including mammals' has a merit that the protein produced is substantially or nearly equivalent to physicochemical properties of the protein obtained as a natural product. For the microbe hosts, for example, *Escherichia coli*, *Bacillus* species, *Streptomyces* species, and yeasts can be arbitrarily used. Examples of the mammalian hosts are epithelial cell, interstitial cell and hemopoietic cell of human, monkey, mouse and hamster origins including 3T3-Swiss albino cell, ATCC CCL-92, C127I cell, ATCC CRL-1616, CHO-K1 cell, ATCC CCL-61, CV-1 cell, ATCC CCL-70, COS-1 cell, ATCC CRL-1650, HeLa cell, ATCC CCL-2, MOP-8 cell, ATCC CRL-

1709, and their mutant strains. To introduce the DNA of this invention into such hosts, one can employ conventional methods, for example, DEAE-dextran method, calcium phosphate transfection method, electroporation method, lipofection method, microinjection method, and viral infection method using retrovirus, adenovirus, herpesvirus and vaccinia virus. To select clones capable of producing the protein among the resulting transformants, the transformants are cultivated in culture media, followed by selection of clones where production of the protein was observed. The above-mentioned recombinant DNA techniques are detailed in, for example, *Jikken-Igaku-Bessatsu, Saibo-Kogaku Handbook* (The handbook for the cell engineering), edited by Toshio KUROKI, Masaru TANIGUCHI and Mitsuo OSHIMURA, published by Yodosha. Co., Ltd., Tokyo, Japan (1992), and *Jikken-Igaku-Bessatsu, Biomanual Series 3, Idenshi-Cloning-Jikken-Ho* (The experimental methods for the gene cloning), edited by Takashi YOKOTA and Kenichi ARAI, published by Yodosha Co., Ltd., Tokyo, Japan (1993).

In this field, once a desired DNA is obtained as described above, then the DNA can be conventionally introduced into animals or plants to establish "transgenic animals" or "transgenic plants". The transgenic animals and plants introduced with the DNA of this invention are also included by the DNA of this invention. The following outlines a procedure for establishing transgenic animals. At first, the DNA of this invention can be introduced into oosperms or embryonic stem cells by using microinjection method, electroporation method or infections with recombinant virus containing the DNA of this invention. Subsequently, thus-obtained cells introduced with the

present DNA can be grafted into uterine tubes or uterus of para-pregnant female animals. Thereafter, from the newborns delivered spontaneously or by caesarean, the transgenic animals introduced with the present DNA can be selected by hybridization method, PCR method, etc. The DNA to be introduced for the establishment of transgenic animals can comprise not only a nucleotide sequence for the present hedgehog protein but also other sequences for promoters or enhancers suitable for regulating the gene expression in desired tissue- and/or stimulation-specific manner and/or further other sequences for signal peptides. Thus, the transgenic animals introduced with the DNA of this invention can be obtained. Techniques for transgenic animals are detailed in a publication such as *Jikken-Igaku-Bessatsu, Shin-Idenshikogaku-Handbook* (The Handbook for Genetic Engineering), edited by Masami MURAMATSU, Hiroto OKAYAMA, and Tadashi Yamamoto, published by Yodosha Co., Ltd., Tokyo, Japan (1996), pp.269-283.

The present hedgehog protein can be prepared by the process of this invention comprising the steps of allowing to express a DNA encoding the hedgehog protein and collecting the generated hedgehog protein. The DNA expression step can include a step of culturing the above-mentioned transformants introduced with the DNA of this invention, encoding the hedgehog protein. The media used to culture the transformant can be selected from conventional ones depending on the types of the transformants to be used, and they are usually composed of, as a base, a bufferized water and, as additives, inorganic ions such as sodium ion, potassium ion, calcium ion, phosphoric ion and chloric ion; microelements, carbon sources, nitrogen sources, amino acids and

vitamins which meet to the metabolism of particular hosts; and, optionally, sera, hormones, cell growth factors and cell adhesion factors. Examples of the carbon sources are saccharides including glucose, fructose, sucrose, starches, and partial hydrolyzates of starches, and examples of the nitrogen sources are nitrogen-containing inorganic and organic substances including ammonia, ammonium ions, urea, nitric ions, peptone, and yeast extracts.

Examples of the culture media are as follows: those for microbe hosts such as L broth medium, T broth medium, TY broth medium, nutrient broth medium, YM broth medium, and potato-dextrose medium; and those for animal hosts such as 199 medium, DMEM medium, Ham's F12 medium, IMDM medium, MCDB104 medium, MCDB153 medium, MEM medium, RD medium, RITC80-7 medium, RPMI-1630 medium, RPMI-1640 medium, WJJC404 medium. To the culture media, the transformant can be inoculated in a cell density of  $1 \times 10^4$ - $1 \times 10^7$  cells/ml, preferably,  $1 \times 10^5$ - $1 \times 10^6$  cells/ml, and cultured under conditions suitable for the hosts, if necessary, while the culture media are replaced with fresh preparations. In particular, when using the hosts of microbe origins, the culture can be carried out at a temperature of 25-65°C and a pH of 5-8 under aerobic conditions such as agitation-aeration for 1-10 days. When using the hosts of animal origins, the culture can be carried out at a temperature of about 37°C for one day to one week, preferably, two to four days by suspension- or monolayer culture. Thus cultures containing the present hedgehog protein are obtained. The content of the present hedgehog protein in the cultures, which may differ depending on the types of the transformants and culture conditions, is usually one microgram



to 100 mg per liter.

Furthermore, in the process for preparing the hedgehog protein of this invention, the DNA expression step can include a step of culturing cells which express the hedgehog protein, for example, established human cell lines ARH-77 cell, ATCC CRL-1621, K-562 cell, ATCC CCL-243, and KU-812 cell, described by K. Kishi et al., in *Leukemia Research*, Vol.9, pp.381-390 (1985). By culturing such cells in culture media suitable for respective cells, for example, 199 medium, DMEM medium, Ham's F12 medium, IMDM medium, MCDB104 medium, MCDB153 medium, MEM medium, RD medium, RITC80-7 medium, RPMI-1630 medium, RPMI-1640 medium, and WJJC404 medium similarly as in culturing of the transformants using animal host cells as mentioned above, then the culture containing the present hedgehog protein can be obtained. The content of the present protein in the cultures, which may differ depending on the types of the cells and culture conditions, is usually one nanogram to one milligram per liter.

The culture products obtained in these manners can be first subjected to ultrasonication, cell-lytic enzyme and/or detergent to disrupt cells, if necessary, the present hedgehog protein can be separated from the cells or cell debris by filtration and centrifugation, followed by purification. In the purification, the culture products which have been separated from cells or cell debris can be subjected to conventional methods used to purify biologically-active proteins, for example, salting-out, dialysis, filtration, concentration, fractional precipitation, ion-exchange chromatography, gel filtration chromatography, adsorption chromatography, isoelectric focusing chromatography, hydrophobic chromatography, reversed phase

chromatography, affinity chromatography, gel electrophoresis and isoelectric focusing gel electrophoresis which are used in combination, if necessary. The purified preparation of the present hedgehog protein can be concentrated and lyophilized into a liquid or solid form to meet to its final use. Immunoaffinity chromatographies using the monoclonal antibody described below do yield a high-purity preparation of the hedgehog protein with minimized costs and labors.

In the process of this invention for producing the hedgehog protein, the DNA expression step can also include a step of feeding or planting the transgenic animals or plants obtained by introducing the DNA which encodes the present hedgehog protein to animals other than humans or plants. After feeding or planting occasionally with desired stimuli, desired tissues, organs, bloods, milks, and/or body fluids of the resultants can be collected and subjected to the steps for purifying the hedgehog protein of this invention as mentioned above to obtain the present protein.

The monoclonal antibody of this invention includes the monoclonal antibodies in general which recognize the hedgehog protein of this invention, independently of its origins, sources, and classes. The monoclonal antibody of this invention can be obtained by using as an antigen the present hedgehog protein, other conventional hedgehog protein or antigenic fragment thereof, and more particularly, by preparing hybridoma cells of derived from an infinitely-proliferative of a mammal and an antibody-producing cell of a mammal that has been immunized with such an antigen, selecting clones of hybridoma capable of producing the monoclonal antibody of this invention, and

culturing the clones *in vitro* or *in vivo*.

Proteins feasible as the antigens can be obtained through culturing of transformants introduced with a DNA encoding at least a partial amino acid sequence of SEQ ID NO:1, and the proteins are usually used after completely or partially purified. The antigenic fragments can be obtained by chemically or enzymatically digesting the completely or partially purified proteins or by chemical synthesis based on the amino acid sequence of SEQ ID NO:1, 2, or 3. Alternatively, the antigens can be obtained by using these techniques based on known hedgehog genes or proteins. Human Sonic hedgehog is useful as such known hedgehog.

Immunization of animals is conducted in conventional manner. For example, the antigens as described above can be injected alone or together with appropriate adjuvants into mammals through an intravenous, intradermal, subcutaneous or intraperitoneal route, and then the mammals can be fed for a prescribed time period. There is no limitation in types of the mammals, therefore any mammals can be used regardless of their types, sizes and genders, as far as one can obtain desired antigen-producing cells therefrom. Rodents such as rats, mice and hamsters are generally used, and among these the most desirable mammal can be chosen in respect to their compatibility with the infinitely-proliferative cells mentioned below. The dose of the antigen is generally set to about five to 500 µg/animal in total, which can be divided into two to five times inoculations with intervals of about one to two weeks, depending on the types and sizes of the mammals to be used. Three to five days after the final inoculation, the spleens are extracted and

dispersed to obtain splenocytes as antibody-producing cells.

The antibody-producing cells obtained in this way can be then fused with infinitely-proliferative cells of mammalian origin to obtain cell-fusion products containing the objective hybridoma. Examples of the infinitely-proliferative cells usually used in this invention are cell lines of mouse myeloma origin such as P3/NSI/1-Ag4-1 cell, ATCC TIB-18; P3X63Ag8 cell, ATCC TIB-9; SP2/O-Ag14 cell, ATCC CRL-1581; and mutant strains thereof. The cell-fusion can be conducted in conventional manner using an electric pulse or a cell-fusion accelerator such as polyethylene glycol and Sendai virus. For example, the antibody-producing cells and the infinitely-proliferative cells of mammalian origin are co-suspended to give a ratio of about 1:1 to 1:10 in a cell fusion medium with such an accelerator and incubated at about 30 to 40°C for about one to five minutes. Although conventional media such as minimum essential medium (MEM), RPMI-1640 medium, and Iscove's modified Dulbecco's medium are feasible as cell fusion media, it is desirable to remove the serum in media, such as bovine serum, prior to their use.

To select the objective hybridomas, the cell-fusion products thus obtained can be transferred to an appropriate selection medium, such as HAT medium, and the hybridomas are cultured at about 30 to 40°C for 3 days to 3 weeks till the cells other than the hybridomas die. The hybridoma cells can be then cultured in usual manner and antibodies secreted in the medium can be tested for reactivity with the hedgehog protein of this invention. Such tests can be conducted in conventional manner directed to detection of antibodies in general, for example, enzyme-immunoassays, radioimmunoassays and bioassays, which are

detailed in *Tan-Clone-Kotai-Jikken-Manual* (Experimental Manual for Monoclonal Antibody), edited by Sakuji TOYAMA and Tamie ANDO, published by Kodansha Scientific, Ltd., Tokyo, Japan (1991), pp.105-152. The hybridomas which recognize the present protein can be immediately cloned by the limiting dilution method, thus obtaining the singly cloned hybridomas according to this invention.

The monoclonal antibody of this invention can be obtained by culturing such hybridomas *in vitro* or *in vivo*. In culturing the hybridomas, conventional methods for culturing mammalian cells can be employed. More particularly, the monoclonal antibody can be collected from culture products in case of culturing *in vitro* in nutrient media, while the monoclonal antibody can be collected from the ascites and/or bloods of the animals in case of transplanting in non-human warm-blooded animals or culturing *in vivo*. To collect the monoclonal antibody from cultures, ascites and blood, conventional methods for purifying antibodies can be arbitrarily used. Particular methods are, for example, salting-out, dialysis, filtration, concentration, fractional precipitation, ion-exchange chromatography, gel filtration chromatography, adsorption chromatography, isoelectric focusing chromatography, hydrophobic chromatography, reversed phase chromatography, affinity chromatography, gel electrophoresis and isoelectric focusing gel electrophoresis which can be used in combination if necessary. The purified preparations of the monoclonal antibody can then be concentrated and dehydrated into liquids or solids to meet to their final use.

The monoclonal antibody of this invention is

particularly useful in immunoaffinity chromatographies for purifying the present hedgehog protein. The method for purification usually comprises the steps of allowing the monoclonal antibody to contact with a mixture of the present protein and contaminants to adsorb the present hedgehog protein on the monoclonal antibody, and desorbing the protein from the antibody; these steps are usually conducted in aqueous systems. The monoclonal antibody of this invention can be used after being immobilized on gels of water-insoluble carriers and packed into columns. For example, the cultures of the transformants or their partially purified preparations are charged to such columns and run, resulting in that the hedgehog protein is substantially-selectively adsorbed by the monoclonal antibody on such carriers. The adsorbed protein can be easily desorbed by altering the hydrogen-ion concentration around the monoclonal antibody. For example, the desorption for eluting the protein is usually conducted under acidic conditions, preferably, pH 2-3 when using the monoclonal antibody belonging to immunoglobulin G (IgG), or alkaline conditions, preferably, pH 10-11 when using the monoclonal antibody belonging to immunoglobulin M (IgM). The present method can yield a high-purity preparation of the present hedgehog protein with minimized costs and labors.

The monoclonal antibody of this invention additionally has wide uses required to detect the present hedgehog protein. The use of the monoclonal antibody in label-immunoassays such as radioimmunoassays, enzyme-immunoassays, and fluorescent-immunoassays can make more rapidly and accurately detect the present hedgehog protein in samples qualitatively or quantitatively. In these immunoassays, the present monoclonal

antibody can be used after being labelled with radioactive substances, enzymes, and/or fluorescent substances. The label-immunoassays have a merit that they can analyze more numerous samples at a time and more accurately than bioassays. Thus the detection method of this invention is significantly useful for quality controls of the present protein during processes of the production or the products, as well as for diagnoses of diseases by detecting the present hedgehog protein. This invention does not basically relate to the techniques for labelling monoclonal antibodies or label-assays, so that it does not describe them in detail. Such techniques are detailed in a publication such as *Enzyme immunoassay*, edited by P. Tijssen, translated by Eiji ISHIKAWA, published by Tokyo-Kagaku-Dojin, Tokyo, Japan (1989), pp.196-348.

The DNA of this invention, which encodes the present hedgehog protein, is also useful in "gene therapies". Particularly, in usual gene therapies, the DNA of this invention can be first inserted into a vector derived from virus such as retrovirus, adenovirus or adeno-associated virus, alternatively, embedded in either cationic- or membrane fusible-liposomes. Subsequently, the inserted or embedded DNA can be directly injected into patients with the hedgehog protein susceptible diseases, alternatively, introduced *in vitro* into lymphocytes, which have been collected from the patients, and self-implanted to the patients. Thus, the DNA of this invention exhibits a remarkable efficacy in gene therapies for diseases being susceptible to human Desert hedgehog protein. General procedures for gene therapies are detailed in *Jikken-Igaku-Bessatsu, Biomanual UP Series, Idenshichiryō-no-Kisogijutsu* (Basic

techniques for the gene therapy), edited by Takashi SHIMADA, Izumi SAITO, and Keiya OZAWA, published by Yodosha Co., Ltd., Tokyo, Japan (1996).

The following Examples describe in detail the way of practicing this invention. The hedgehog protein of this invention, the DNA encoding the hedgehog protein, and the process for producing the hedgehog protein are explained by Examples 1 to 3, the monoclonal antibody of this invention and process for preparing the antibody are explained by Example 4, and the method for detecting the hedgehog protein using the monoclonal antibody of this invention is explained by Examples 5 and 6. The following Examples can be diversified by the technical level in this field. In view of this, this invention should not be restricted to the Examples:

#### Example 1

##### Preparation of DNA

##### Example 1-1(a)

##### Preparation of total RNA

ARH-77 cells, ATCC CRL-1621, an established cell line derived from human plasma cell leukemia, were suspended in RPMI-1640 medium supplemented with 10%(v/v) fetal bovine serum and proliferated in usual manner at 37°C in a 5%(v/v) CO<sub>2</sub> incubator while scaling up the culture. After the cell density reached a desired level, the cells were collected. The cells were suspended in micro-centrifugal tubes with phosphate-buffered saline (hereinafter, abbreviated as "PBS") and centrifuged, and the supernatants were discarded; these treatments were repeated three times. Then the cells were placed in fresh micro-centrifugal tubes in an amount of 5×10<sup>6</sup> cells/tube, and



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"ULTRASPEC™ RNA", a total RNA isolation reagent commercialized by BIOTECHX Laboratories, Inc., Houston, Texas, USA, was added to the tubes in a volume of 1.0 ml/tube before the cells were suspended. The suspensions were allowed to stand in ice-chilling conditions for 5 minutes, mixed with 1.2 ml/tube of a mixture of chloroform/"ULTRASPEC™ RNA" (1/5 by volume), shaken for 15 seconds, and allowed to stand in ice-chilling conditions for 5 minutes. Upper phase in the tubes formed by centrifugation was collected, mixed with the equal volume of 2-propanol, and allowed to stand in ice-chilling conditions for five minutes. The mixture was centrifuged, and the supernatant was discarded. The formed precipitate was washed twice with 75%(v/v) aqueous ethanol, dried up in vacuo, and dissolved in sterile distilled water, resulting in obtaining an aqueous solution containing total RNAs of ARH-77 cells. A small portion of the solution was examined for the absorbance at 260 nm to calculate an RNA content.

#### Example 1-1(b)

##### Preparation of first strand cDNA

Based on the nucleotide sequence of a mouse Desert hedgehog gene registered in "GenBank®", a nucleic acid database by National Institute of Health, USA, under the accession number "X76292", an oligonucleotide with the nucleotide sequence of 5'-GCCAGGGTGTGAGCAACAGT-3' (SEQ ID NO:12) was prepared in usual manner. In a micro-reaction tube, 2.5 pmol of the oligonucleotide and one microgram of total RNAs prepared by the method in Example 1-1(a) were placed, and sterile distilled water was added to the mixture to give a final volume of 15.5 µl. After the tube was allowed to stand at 70°C for ten minutes and

under ice-chilling conditions for one minute, to the tube 2.5 µl of 10 × PCR buffer, 2.5 µl of 25 mM MgCl<sub>2</sub>, 1.0 µl of 10 mM dNTP mix, and 2.5 µl of 0.1 M DTT were added in this order. The tube was allowed to stand at 42°C for one minute. First strand cDNAs was synthesized by adding to the tube one microliter of "SUPERScript II RT", a reagent of reverse transcriptase commercialized by GIBCO BRL, Life Technologies, Inc., Rockville, Maryland, USA, and incubating the tube at 42°C for 50 minutes. After the mixture was heated to terminate the reaction at 70°C for 15 minutes and cooled to 37°C, the RNAs were degraded by incubating with admixed one microliter of RNase at 37°C for 30 minutes. Thereafter, from the reaction mixture, an aqueous solution containing purified first strand cDNAs in a volume of 50 µl was obtained by mixing with 120 µl of 6 M NaI and treating with "GlassMAX™", a DNA isolation matrix commercialized by GIBCO BRL, Life Technologies, Inc., Rockville, Maryland, USA, in accordance with the accompanying instructions.

#### Example 1-1(c)

#### Preparation of DNA fragment encoding the hedgehog protein and recombinant DNA

Ten-microliter portion of a solution of first strand cDNAs, obtained by the method in Example 1-1(b), was sampled in a micro-reaction tube and manipulated with "5' RACE SYSTEM, VERSION 2.0", a kit for a modified PCR method of 5' RACE, commercialized by GIBCO BRL Life Technologies, Inc., Rockville, Maryland, USA, in accordance with the accompanying instructions to add a poly(C)-tail to each of the 5'-termini of the cDNAs and amplify DNA fragments for the 5'-terminal regions. The sense primer used was "anchor primer" in the kit, and the antisense

primer used was the oligonucleotide in Example 1-1(b). The thermal controls were as follows: an incubation at 94°C for one minute; 35 cycles of incubations at 94°C for one minute, at 55°C for one minute, and at 72°C for one minute; and an incubation at 72°C for 10 minutes. The reaction volume was set to 50 µl.

A DNA fragment which encodes a Desert hedgehog protein of human origin was obtained by PCR using the above reaction mixture as a template under conditions as follows. Sense and antisense primers for this PCR were obtained in usual manner based on the nucleotide sequence of a Desert hedgehog protein of mouse origin, which is reported by Y. Echelard et al. and registered in "GenBank®", a nucleic acid database established by National Institute of Health, USA, under the accession number "X76292"; they had respective nucleotide sequences of 5'-TGTGCTGCTTGGCACTCTTG-3' (SEQ ID NO:13) and 5'-CCGTGGCATTTCCTCGGAAAG-3' (SEQ ID NO:14). Two microliters of 100-folds dilution of the reaction mixture of the above 5' RACE was placed in a fresh micro-reaction tube, then to which 3 µl of 10 × PCR buffer, 1.8 µl of 25 mM MgCl<sub>2</sub>, 0.6 µl of 10 mM dNTP mix, appropriate amounts of the sense and antisense primers, and sterile distilled water were added to give a final volume of 30 µl. After 0.3 µl of 5 units/µl Taq DNA polymerase was added to the tube, the mixture was subjected to an incubation at 94°C for three minutes, 35 cycles of incubations at 94°C for one minute, at 55°C for one minute, and at 72°C for one minute, and finally an incubation at 72°C for 10 minutes, to effect PCR. The PCR products were subjected to 2%(w/v) agarose gel electrophoresis. A gel portion containing an about 600 bp-DNA band, stained with ethidium bromide, was excised and treated with "SUPREC™-01", a

DNA purification tube commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, to obtain 20 µl aqueous solution containing a DNA fragment.

A portion of the DNA fragment solution was sampled and manipulated with "pCR-SCRIPT SK(+) CLONING KIT", a DNA cloning kit commercialized by Stratagene Cloning Systems, California, USA, in accordance with the accompanying instructions to ligate the DNA fragment with "pCR-SCRIPT SK(+)", the plasmid vector in the kit. After the ligation, a portion of the reaction mixture was introduced by conventional transformation method into competent cells of *Escherichia coli* "JM101" strain, commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, which were then inoculated on plates of L agar medium containing 50 µg/ml ampicillin and cultured at 37°C under standing conditions overnight. Some of the colonies formed were respectively suspended in 10 µl aliquotes of sterile distilled water. PCRs were conducted under the same conditions as described above in this Example except for using the suspensions as respective templates. Colonies which gave an about 600 bp-DNA on agarose gel electrophoresis were respectively inoculated to aliquotes of L broth medium containing 50 µg/ml ampicillin and cultured at 37°C under shaking conditions overnight. From the resulting cultures, recombinant DNAs were collected by conventional alkali-sodium dodecyl sulfate (hereinafter, sodium dodesyl sulfate is abbreviated as "SDS") method. The recombinant DNAs were sequenced by dideoxy method. The DNA fragment in the recombinant DNAs contained the nucleotide sequence of SEQ ID NO:7.

Studying homology between thus-determined nucleotide sequence and other known nucleotide sequences, the nucleotide

sequence determined in this Example exhibited a significant homology of about 89% to the nucleotide sequence of a mouse Desert hedgehog gene, registered in "GenBank®", a nucleic acid database by National Institute of Health, USA, under the accession number "X76292". This indicates that the DNA fragment of this Example encodes a human Desert hedgehog protein. The recombinant DNA obtained in this Example was named "pHuDHH/#20". In addition, the nucleotide sequence of SEQ ID NO:7, determined in this Example, was compared with the informations on structures and functions of known hedgehog proteins as described by M. Hammerschmidt et al., in *Trends in Genetics*, Vol.13, pp.14-21 (1997), leading to a conclusion that the sequence of nucleotides 19-546 of SEQ ID NO:7 encodes a mature form of a human Desert hedgehog protein and that the mature form of the protein can contain the amino acid sequence shown along with this nucleotide sequence, which is also shown in SEQ ID NO:1.

#### Example 1-2

##### Preparation of DNA fragment and recombinant DNA encoding the hedgehog protein

A recombinant DNA "pHuDHH/#20", obtained by the method in Example 1-1(c), was cleaved with restriction enzymes *EcoRI* and *NotI* and subjected to 2%(w/v) agarose gel electrophoresis. From the agarose gel, a gel portion containing an about 600 bp-DNA band stained with ethidium bromide was excised and treated with "SUPREC™-01", a DNA purification tube commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, to collect and purify a DNA fragment. The purified DNA fragment was labelled with a radioisotope <sup>32</sup>P by using a DNA labelling kit, "MEGAPRIME™ DNA LABELLING SYSTEMS", commercialized by Amersham Corp., Div.

Amersham International, Arlington Heights, USA, and manipulating in accordance with the accompanying instructions. Human fetal brain cDNA library constructed with a type of bacteriophage  $\lambda$ gt11 as a vector, commercialized by CLONTECH Laboratories, Inc., Palo Alto, California, USA, was screened by using the  $^{32}\text{P}$ -labelled DNA fragment as a probe. Cells of *Escherichia coli* strain "NM514" were infected with the cDNA library in usual manner, inoculated on plates of L agar medium, and cultured at  $37^\circ\text{C}$  for 6-18 hours to form plaques. The formed plaques were transferred in usual manner to nylon membranes, which were then subjected to alkali-denaturalization, neutralization, and air-drying in usual manner. The air-dried membranes were immersed in a pre-hybridization solution, which contained  $6 \times \text{SSC}$ ,  $5 \times \text{Denhardt's solution}$ ,  $0.5\%(\text{w/v})$  SDS,  $50\%(\text{v/v})$  formamide, and  $100 \mu\text{g/ml}$  denatured salmon sperm DNA, at  $42^\circ\text{C}$  for 1-2 hours, and subsequently immersed in a fresh pre-hybridization solution with an appropriate amount of the  $^{32}\text{P}$ -labelled DNA fragment added as a probe and incubated at  $42^\circ\text{C}$  for 16-20 hours to effect hybridization. After the hybridization, the membranes were washed with  $2 \times \text{SSC}$  containing  $0.1\%(\text{w/v})$  SDS at ambient temperature for 15 minutes and further washed with  $0.2 \times \text{SSC}$  containing  $0.1\%(\text{w/v})$  SDS at a temperature moderately increasing from  $37^\circ\text{C}$  to  $65^\circ\text{C}$  until background radioactivity was adequately reduced. Thereafter the membranes were subjected to autoradiography. From a plaque which gave a positive signal, a phage clone was collected and amplified in usual manner, and from the amplified phage a DNA clone was collected. The DNA clone was sequenced by dideoxy method using primers prepared based on the vector's nucleotide sequence. The DNA clone contained a partial nucleotide sequence as shown with

5'-GTATCCATGGCTCTCCTG-3' (SEQ ID NO:15). Compared with other known nucleotide sequences, the partial nucleotide sequence had a significant homology to a partial nucleotide sequence, containing translation initiation site, of a mouse Desert hedgehog gene registered in "GenBank®", a nucleic acid database by National Institute of Health, USA, under the accession number "X76292".

As sense and antisense primers for PCR, oligonucleotides with respective nucleotide sequences of 5'-GCCTCGAGGTATCCATGGCTCTCCTG-3' (SEQ ID NO:16), which contains the above-determined partial nucleotide sequence, and 5'-GCGCGGCCGCTCAGCCGCCCGCCGGAC-3' (SEQ ID NO:17), which is complementary to the sequence of nucleotides 532-548 of SEQ ID NO:7, were prepared in usual manner. As a template one microliter portion of cDNAs solution, obtained by the methods in Examples 1-1(a) and 1-1(b), was placed in a micro-reaction tube, then to which 3  $\mu$ l of 10  $\times$  PCR buffer, 1.8  $\mu$ l of 25 mM  $MgCl_2$ , 0.6  $\mu$ l of 10 mM dNTP mix, appropriate amounts of the sense and antisense primers, and sterile distilled water were added to give a final volume of 30  $\mu$ l. After 0.3  $\mu$ l of 5 units/ $\mu$ l Taq DNA polymerase was added to the tube, the mixture was subjected to an incubation at 94°C for three minutes, 35 cycles of incubations at 94°C for one minute, at 55°C for one minute, and at 72°C one minute, and finally an incubation at 72°C for 10 minutes, to effect PCR. The PCR products were subjected to 2%(w/v) agarose gel electrophoresis. From the gel, a gel portion containing an about 600 bp-DNA band, stained with ethidium bromide, was excised and treated with "SUPREC™-01", a DNA purification tube commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, to obtain

20 µl aqueous solution containing a DNA fragment.

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A small portion of the DNA fragment solution was sampled and manipulated with "pT7BLUE CLONING KIT", a DNA cloning kit commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, in accordance with the accompanying instructions to ligate the DNA fragment with "pT7BLUE", the plasmid vecotor in the kit. After the ligation, a portion of the reaction mixture was introduced by usual transformation method into competent cells of *Escherichia coli* strain "JM101", commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, which were then inoculated on plates of L agar medium containing 50 µg/ml ampicillin and cultured at 37°C under standing conditions overnight. The formed colonies were respectively suspended in 10 µl aliquotes of sterile distilled water. Except for using the suspensions as respective templates, PCRs were conducted under the same conditions as in Example 1-1(c). Colonies which gave an about 600 bp-DNA band on agarose gel electrophoresis were respectively inoculated to aliquotes of L broth medium containing 50 µg/ml ampicillin and cultured at 37°C under shaking conditions overnight. From the resulting cultures recombinant DNAs were collected by conventional alkali-SDS method. The recombinant DNAs were sequenced by dideoxy method. The DNA fragment in the recombinant DNAs contained the nucleotide sequence of SEQ ID NO:8, which can encode the amino acid sequence shown along with the nucleotide sequence.

The nucletotide sequence of SEQ ID NO:8 was compared with the nucleotide sequence of SEQ ID NO:7, determined in Example 1-1. The sequence of nucleotides 1-548 of SEQ ID NO:7 completely matched with the sequence of nucleotides 55-602 of SEQ ID NO:8. The results of this comparison and the comparison with



the above-mentioned nucleotide sequence of a mouse Desert hedgehog gene revealed that: the nucleotide sequence of SEQ ID NO:8 encodes N-terminal region of a precursor form of a human Desert hedgehog protein; the sequence of nucleotides 7-72 of SEQ ID NO:8 encodes a signal peptide in a precursor form of the hedgehog protein; and the sequence of nucleotides 73-600 of SEQ ID NO:8 encodes a mature form of the hedgehog protein which contains the amino acid sequence of SEQ ID NO:1.

#### Example 1-3

##### Preparation of DNA fragment and recombinant DNA encoding the hedgehog protein

As sense and antisense primers for PCR, oligonucleotides with respective nucleotide sequences of 5'-CGTGTCGGTCAAAGCTGATA-3' (SEQ ID NO:18) and 5'-ATGCATTCCAGTCGGCTGGA-3' (SEQ ID NO:19) were prepared in usual manner; the former sequence was identical to the sequence of nucleotides 501-520 of SEQ ID NO:7, and the latter sequence was based on the nucleotide sequence registered in "GenBank®", a nucleic acid database by National Institute of Health, USA, under the accession number "AA064660", which is of a human cDNA fragment similar to a 3'-terminal sequence for a mouse Desert hedgehog protein in a precursor form. As a template one microliter portion of cDNAs solution, obtained by the methods in Examples 1-1(a) and 1-1(b), was placed in a micro-reaction tube, then to which 3 µl of 10 × PCR buffer, 1.8 µl of 25 mM MgCl<sub>2</sub>, 0.6 µl of 10 mM dNTP mix, appropriate amounts of the above sense and antisense primers, and sterile distilled water were added to give a final volume of 30 µl. After 0.3 µl of 5 units/µl Taq DNA polymerase was added to the tube, the mixture was subjected to

an incubation at 94°C for three minutes, 35 cycles of incubations at 94°C for one minute, at 55°C for one minute, and at 72°C one minute, and finally an incubation at 72°C for 10 minutes to, effect PCR. The PCR products were subjected to 2%(w/v) agarose gel electrophoresis. From the gel, a gel portion containing an about 600 bp-DNA band, stained with ethidium bromide, was excised and treated with "SUPREC™-01", a DNA purification tube commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, to obtain 20 µl aqueous solution containing a DNA fragment.

A portion of the DNA fragment solution was sampled and manipulated with "pT7BLUE CLONING KIT", a DNA cloning kit commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, in accordance with the accompanying instructions to ligate the DNA fragment with "pT7BLUE", the plasmid vector in the kit. After the ligation, a portion of the reaction mixture was introduced by usual transformation method into competent cells of *Escherichia coli* strain "JM101", commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, which were then inoculated on plates of L agar medium containing 50 µg/ml ampicillin and cultured at 37°C under standing conditions overnight. The formed colonies were respectively suspended in 10 µl aliquotes of sterile distilled water. Except for using the suspensions as respective templates and using the sense and antisense primers in this Example, PCRs were conducted under the same conditions as in Example 1-1(c). Colonies which gave an about 600 bp-DNA band on agarose gel electrophoresis were respectively inoculated to aliquotes of L broth medium containing 50 µg/ml ampicillin and cultured at 37°C under shaking conditions overnight. From the resulting cultures recombinant DNAs were collected by conventional alkali-SDS

method. The recombinant DNAs were sequenced by dideoxy method. The DNA fragment in the recombinant DNAs contained the nucleotide sequence of SEQ ID NO:9, which can encode the amino acid sequence shown along with the nucleotide sequence.

The nucleotide sequence of SEQ ID NO:9 was compared with the nucleotide sequence of SEQ ID NO:7, determined in Example 1-1. The sequence of nucleotides 501-548 of SEQ ID NO:7 completely matched with the sequence of nucleotides 1-48 of SEQ ID NO:9. In addition, the nucleotide sequence of SEQ ID NO:9 was compared with the informations on structure and function of known hedgehog proteins as described by M. Hammerschmidt et al., in *Trends in Genetics*, Vol.13, pp.14-21 (1997), revealing that the nucleotide sequence of SEQ ID NO:9 partially encodes a C-terminal region of a precursor form of a human Desert hedgehog protein.

#### Example 1-4

#### Preparation of DNA fragment and recombinant DNA encoding the hedgehog protein

Total RNAs in a weight of 1.5 µg, obtained by the method in Example 1-1(a), was placed in a micro-reaction tube, to which 2 µl of 5 × reverse transcriptase buffer, 2 µl of DTT, one microliter of 10 mM dNTP mix, an appropriate amount of an oligonucleotide as an adaptor primer for a modified PCR method of 3' RACE with the nucleotide sequence of 5'-AAGGATCCGTCGACAAGCTTAATACGACGAATTCTGGAG(T)<sub>17</sub>-3' (SEQ ID NO:20) prepared in usual manner, and sterile distilled water were added to give a final volume of 29 µl. After admixed with one microliter of "SuperScript™ II RT", a reagent of reverse transcriptase commercialized by GIBCO BRL Life Technologies Inc., Rockville, Maryland, USA, the tube was allowed to stand at 37°C

for about 1.5 hours to effect a reaction of synthesizing cDNAs to poly(A)'RNAs 3'-terminal regions. Except for using a portion of the reaction mixture as a template and oligonucleotides with respective nucleotide sequences of 5'-GGCTTCGACTGGGTCTACTA-3' (SEQ ID NO:21) as a sense primer and 5'-AAGGATCCGTCGACAAG-3' (SEQ ID NO:22) as an antisense primer prepared in usual manner, a first step PCR was conducted under the same conditions as in Example 1-3; the sequence of the sense primer was identical to the sequence of nucleotides 460-479 of SEQ ID NO:7, and that of the antisense primer was based on the above adaptor primer. After the first step PCR, the reaction mixture was diluted with an appropriate amount of sterile distilled water. Except for using the dilution as a template and oligonucleotides with the respective nucleotide sequences of 5'-ATGCGCTTCGGCCAGCG-3' (SEQ ID NO:23) as a sense primer and 5'-GACAAGCTTAATACGAC-3' (SEQ ID NO:24) as an antisense primer, a second step PCR was conducted under the same conditions as in Example 1-3; the sequence of the sense primer was identical to the sequence of nucleotides 369-385 of SEQ ID NO:9, and that of the antisense primer was based on the nucleotide sequence of the above adaptor primer. After the second step PCR, the reaction mixture was diluted with an appropriate amount of sterile distilled water. Except for using the dilution as a template and oligonucleotides with the respective nucleotide sequences of 5'-GTTTCGCGCCGCTCACCG-3' (SEQ ID NO:25) as a sense primer and 5'-TACGACGAATTCTGGAGT-3' (SEQ ID NO:26) as an antisense primer, a third step PCR was conducted under the same conditions as in Example 1-3; the sequence of the sense primer was identical to the sequence of nucleotides 424-440 of SEQ ID NO:9, and that of the antisense primer was based on the

nucleotide sequence of the above adaptor primer. The third step PCR products were subjected to 2%(w/v) agarose gel electrophoresis. From the gel, a gel portion containing an about 750 bp-DNA band stained with ethidium bromide was excised and treated with "SUPREC™-01", a DNA purification tube commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, to obtain 20 µl aqueous solution containing a DNA fragment.

A portion of the DNA fragment solution was sampled and manipulated with "pT7BLUE CLONING KIT", a DNA cloning kit commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, in accordance with the accompanying instructions to ligate the DNA fragment with "pT7BLUE", the plasmid vector in the kit. After the ligation, a portion of the reaction mixture was introduced by usual transformation method into competent cells of *Escherichia coli* strain "JM101", commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, which were then inoculated on plates of L agar medium containing 50 µg/ml ampicillin and cultured at 37°C under standing conditions overnight. The formed colonies were respectively suspended in 10 µl aliquotes of sterile distilled water. Except for using the suspensions as respective templates and using the sense and antisense primers in the third step PCR, PCRs were conducted under the same conditions as in Example 1-1(c). Colonies which gave an about 750 bp-DNA band on agarose gel electrophoresis were respectively inoculated to aliquotes of L broth medium containing 50 µg/ml ampicillin and cultured at 37°C under shaking conditions overnight. From the resulting cultures recombinant DNAs were collected by alkali-SDS method. The recombinant DNAs were sequenced by dideoxy method. The DNA fragment in the recombinant DNAs contained the nucleotide

sequence of SEQ ID NO:10, which can encode the amino acid sequence shown along with the nucleotide sequence.

The nucleotide sequence of SEQ ID NO:10 was compared with the nucleotide sequence of SEQ ID NO:9, determined in Example 1-3. The sequence of nucleotides 1-152 of SEQ ID NO:10 completely matched with the sequence of nucleotides 424-575 of SEQ ID NO:9. The results of this comparison and the comparison with the above-mentioned nucleotide sequence of a mouse Desert hedgehog gene revealed that the nucleotide sequence of SEQ ID NO:10 encodes a region containing the C-terminus of a precursor form of a human Desert hedgehog protein.

As described in Examples 1-1 to 1-4, the nucleotide sequences of SEQ ID NOs:7-10, determined in these Examples, were proved to be overlapping nucleotide sequences one another which partially encode a precursor form of a human Desert hedgehog protein; and the precursor protein can be wholly encoded by a DNA containing the nucleotide sequence of SEQ ID NO:6. In addition, these results elucidated that: a human Desert hedgehog protein can be in a precursor form which contains the amino acid sequence of SEQ ID NO:2 or 3 or in a mature form which contains the amino acid sequence of SEQ ID NO:1; such precursor protein can be encoded by a DNA containing the nucleotide sequence of SEQ ID NO:5 or 6, respectively; and such mature protein can be encoded by a DNA containing the nucleotide sequence of SEQ ID NO:4.

#### Example 2

##### Preparation of transformant

Based on the nucleotide sequence determined in Example 1-1(c), which encodes a precursor form of a human Desert hedgehog protein, oligonucleotides with respective nucleotide sequences

of 5'-CCCGGGAATTCATTGCGGGCCGGGCCGGGGGCCG-3' (SEQ ID NO:27) as a sense primer and 5'-ACGATGAATTCTCAGCCGCCCGCCCGGACCGCCA-3' (SEQ ID NO:28) as an antisense primer were prepared in usual manner. PCR was conducted under the same conditions as in Example 1-1(c) except for using the recombinant DNA "pHuDHH/#20" as a template, obtained by the method in Example 1-1(c), and the above sense and antisense primers. An about 600 bp-DNA amplified in this PCR was purified by 2%(w/v) agarose gel electrophoresis and treating with "SUPREC™-01", a DNA purification tube commercialized by Takara Shuzo Co., Ltd., Tokyo, Japan, in accordance with the accompanying instruction, to obtain 20 µl aqueous DNA solution. Two microliters portion of the DNA solution was sampled and subjected to a ligation reaction using T4 DNA ligase with "pCR™II", a plasmid vector for TA cloning commercialized by Invitrogen Corporation, San Diego, USA. A portion of the reaction mixture was introduced by usual transformation method into competent cells of *Escherichia coli* strain "TOP10F'", commercialized by Invitrogen Corporation, San Diego, USA, which were then inoculated on plates of L agar medium containing 50 µg/ml ampicillin and 50 µg/ml 5-bromo-4-chloro-3-indolyl-β-D-galactoside and cultured at 37°C under standing conditions. A white colony formed was inoculated to an aliquote of L broth medium containing 50 µg/ml ampicillin and cultured at 37°C under shaking conditions overnight. From the resulting culture, a recombinant DNA was collected by alkali-SDS method. The recombinant DNA was treated with restriction enzyme *EcoRI* and subjected to 2%(w/v) agarose gel electrophoresis, on which an about 600 bp-DNA was separated, and it was then purified with "SUPREC™-01", a DNA purification tube commercialized by Takara

Shuzo Co., Ltd, Tokyo, Japan.

A portion of the purified DNA solution was sampled and subjected to a ligation reaction in usual manner using T4 DNA ligase with plasmid vector "pGEX-2T", commercialized by Pharmacia Biotech, Inc., Uppsala, Sweden, which had been cleaved with *EcoRI* and dephosphorylated prior to use. A portion of the ligation reaction mixture was introduced by usual transformation method into competent cells prepared by applying the method in *DNA cloning*, Vol.1, edited by D. M. Glover, published by IRL press limited, Oxford, England (1985), pp.109-136, to *Escherichia coli* "BL21" strain, commercialized by Pharmacia Biotech, Inc., Uppsala, Sweden, which were then inoculated to plates of L agar medium containing 50 µg/ml ampicillin and cultured at 37°C under standing conditions overnight. A colony formed was inoculated to an aliquote of L both medium containing 50 µg/ml ampicillin and cultured at 37°C under shaking conditions overnight. From the resulting culture, a recombinant DNA was collected by alkali-SDS method. The recombinant DNA was confirmed by dideoxy method to contain the nucleotide sequence of SEQ ID NO:4, encoding the amino acid sequence of SEQ ID NO:1. The recombinant DNA and the transformant with the recombinant DNA introduced, thus obtained, were named "pHuDHH5'/pGEX-2T/#4-8" and "TAL#4-8/HuDHH", respectively. As shown in FIG.1, in the recombinant DNA "pHuDHH5'/pGEX-2T/#4-8", the DNA with the nucleotide sequence of SEQ ID NO:4 encoding a mature form of a human Desert hedgehog protein and a termination codon were respectively located in the downstream and further downstream of a structural gene of glutathione S-transferase in the same frame with the gene, which was under the regulation of Tac promotor.



### Example 3

#### Production of the hedgehog protein

A transformant "TAL#4-8/HuDHH" obtained by the method in Example 2 was cultured in L broth medium containing 50 µg/ml ampicillin at 37°C under shaking conditions overnight to obtain a seed culture. One milliliter of the seed culture was added to 100 ml of the same medium, freshly prepared in a 500 ml-Erlenmeyer flask, and cultured at 37°C under shaking conditions while the absorbance at 600 nm was monitored. When the absorbance reached a value of 0.5, 0.1 ml of 100 mM isopropylthio-β-D-galactoside was added to the culture. After further cultivation at 37°C for 3.5 hours, the cells were collected from the culture by centrifugation. The cells were washed with PBS, suspended in 5 ml of fresh preparation of PBS, and disrupted with ultrasonication in usual manner. After the cell-disruptant was centrifuged, the formed supernatant was collected.

The supernatant was added to "GLUTATHIONE SEPHAROSE 4B BEADS", a preparation of sepharose beads linked to glutathione, commercialized by Pharmacia Biotech, Inc., Uppsala, Sweden, and incubated at ambient temperature for 30 minutes. After centrifugation of the mixture and discard of the resulting supernatant, the beads were washed twice with PBS. To the beads, an appropriate amount of 50 mM Tris-HCl buffer (pH 7.5) containing 2.5 mM CaCl<sub>2</sub> and 150 mM NaCl was added, and admixed with 10 units of thrombin, commercialized by Ito Ham Co., Ltd., Nishinomiya, Japan, per one milligram of the proteinaceous components. The mixture was incubated at ambient temperature for 16 hours. The mixture was centrifuged to collect a supernatant,

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which was then admixed with an appropriate amount of "ANTITHROMBIN AGAROSE", commercialized by Sigma Chemical Company, St. Louis, Missouri, USA, and centrifuged. The resulting supernatant was added to "HEPARIN AGAROSE", commercialized by Sigma Chemical Company, St. Louis, Missouri, USA, previously equilibrated with equilibration buffer (PBS containing 1.0 mM DTT and 0.2 mM phenylmethanesulfonyl fluoride), and incubated at ambient temperature for 30 minutes. The mixture was admixed with an appropriate amount of equilibration buffer and centrifuged, and the resulting supernatant was discarded. To the remaining components an appropriate amount of 650 mM NaCl was added, and the resulting mixture was centrifuged to collect a supernatant. These treatments, i.e., addition of 650 mM NaCl, centrifugation, and collection of a supernatant, were additionally applied twice to the remaining components, and the supernatants thus obtained were pooled.

A portion of the pooled liquid was subjected to SDS-polyacrylamide gel electrophoresis (hereinafter, abbreviated as "SDS-PAGE") in the presence of a reducing agent, in accordance with the method by U. K. Laemli, in *Nature*, Vol.227, pp.680-685 (1970). As molecular weight markers "SDS-PAGE STANDARDS, LOW RANGE", containing six proteins with distinctive molecular weights of 14,400-97,400 daltons, commercialized by Bio-rad Laboratories Inc., Richmond, USA, was used. Main bands were observed at positions corresponding to molecular weights of about  $22,000 \pm 2,000$  daltons and about  $18,000 \pm 2,000$  daltons. Reference 1 was conducted in the same manner as in this Example except for using *Escherichia coli* "BL21" strain in place of the transformant "TAL#4-8/HuDHH", giving no remarkable band on SDS-PAGE.

Reference 2 was conducted in the same manner as this Example except for using *Escherichia coli* "BL21" strain transformed with the plasmid vector "pGEX-2T" in place of the transformant "TAL#4-8/HuDHH", giving no remarkable band on SDS-PAGE.

The molecular weight of a human Desert hedgehog protein in a mature form which has the amino acid sequence of SEQ ID NO:1 is calculated to be 19,747. According to this Example, the objective protein is usually generated in a form with a peptide as shown by Gly-Ser-Pro-Gly-Ile-His- (SEQ ID NO:29) added to the N-terminus and collected. The molecular weight of a protein that has the amino acid sequence of SEQ ID NO:1 and Gly-Ser-Pro-Gly-Ile-His- (SEQ ID NO:29), which is added to the N-terminus of SEQ ID NO:1, is calculated to be 20,296. These indicate that the protein obtained by the method in this Example which gave a molecular weight of 22,000±2,000 daltons on SDS-PAGE is a type of the present hedgehog protein, containing the amino acid sequence of SEQ ID NO:1. The other protein obtained by the method in this Example, which gave a molecular weight of 18,000±2,000 daltons on SDS-PAGE, is considered to be a degradation product of the hedgehog protein formed during the process of this Example. These results mean that the process of this invention satisfactorily produces a human Desert hedgehog protein of this invention.

#### Example 4

##### Production of monoclonal antibody

##### Example 4-1

##### Preparation of immunogen

##### Example 4-1(a)

##### Preparation of transformant introduced with DNA that encodes

immunogen

A549 cells, ATCC CCL-185, an established cell line derived from a human lung carcinoma, were suspended in RPMI-1640 medium (pH 7.2) supplemented with 10%(v/v) fetal bovine serum and proliferated in usual manner at 37°C in a 5%(v/v) CO<sub>2</sub> incubator while scaling up the culture. After the cell density reached a desired level, proliferated cells were collected. The cells were manipulated with "ULTRASPEC™ RNA", similarly as in Example 1-1(a), to obtain an aqueous solution containing total RNAs of A549 cells. By applying usual RT-PCR method to the total RNAs, a DNA fragment encoding a mature form of a human Sonic hedgehog protein was amplified. As the sense and antisense primers in this RT-PCR, oligonucleotides with respective nucleotide sequences of 5'-CCCGGGAATTCATTGCGGACCGGGCAGGGGGTT-3' (SEQ ID NO:30) and 5'-ACGATGAATTCTCAGCCTCCCGATTGGCCGC-3' (SEQ ID NO:31), prepared in usual manner based on the nucleotide sequence of a human Sonic hedgehog gene, reported by V. Marigo et al. and registered in "GenBank®", a nucleic acid database by National Institute of Health, USA, under the accession number "L38518", were used. The amplified DNA fragment was collected by treating the RT-PCR products with "SUPREC™-01", as in Example 1-1(c). Similarly as in Example 1-1(c), the DNA fragment was ligated with plasmid vector "pCR™II" and introduced into *Escherichia coli* "TOP10F'" strain, the obtained transformant was cultured, and from the resulting culture a recombinant DNA was collected by alkali-SDS method. The recombinant DNA was sequenced by dideoxy method, confirming that it contained the nucleotide sequence of SEQ ID NO:11, encoding a human Sonic hedgehog protein in a mature form.

Similarly as in Example 2, an aliquot of the

recombinant DNA was cleaved with restriction enzyme *EcoRI* to form an about 600 bp-DNA, which was then collected by treating with "SUPREC™-01", ligated with plasmid vector "pGEX-2T", and introduced into *Escherichia coli* "BL21" strain. The obtained transformant was cultured, and from the resulting culture a recombinant DNA was collected by alkali-SDS method. The recombinant DNA was sequenced by dideoxy method, confirming that it contained a DNA with the nucleotide sequence of SEQ ID NO:11 and a termination codon, which are respectively located in the downstream and further downstream of a structural gene of glutathione S-transferase in the same frame under the regulation of Tac promotor. The recombinant DNA and the transformant with the recombinant DNA introduced, thus obtained, were named "pHuSHH/pGEX-2T/#3-1" and "TAL#3-1/HuSHH", respectively.

#### Example 4-1(b)

##### Preparation of immunogen using transformant

Similarly as in Example 3, the transformant "TAL#3-1/HuSHH" obtained by the method in Example 4-1(a) was cultured, the proliferated cells were collected from the culture, and a supernatant of the cell-disruptant was obtained. By applying the methods using "GLUTATHIONE SEPHAROSE 4B BEADS", thrombin, "ANTITHROMBIN AGAROSE" and "HEPARIN AGAROSE" in Example 3 to the supernatant, an aqueous solution containing a protein derived from "TAL#3-1/HuSHH" was obtained, and analyzed by SDS-PAGE; a main band was observed at a position corresponding to a molecular weight of  $22,000 \pm 2,000$ . The molecular weight of a mature form of a human Sonic hedgehog protein which has the amino acid sequence shown along with SEQ ID NO:11 is calculated to be 19,747. According to this Example, the objective protein is

usually generated in a form with a peptide as shown by Gly-Ser-Pro-Gly-Ile-His- (SEQ ID NO:29) added to the N-terminus and collected. These indicate that the protein obtained in this Example is a human Sonic hedgehog protein with a satisfactory purity. Thus, a purified preparation of a human Sonic hedgehog protein as an immunogen was obtained.

#### Example 4-2

##### Preparation of hybridoma

Seven-week-old BALB/c mice were intraperitoneally injected with a purified preparation of a human Sonic hedgehog protein, obtained by the method in Example 4-1(b), in a dose of 100 µg/body together with complete Freund adjuvant in usual manner. Two weeks later, the above injection was repeated, and then the mice were injected with incomplete Freund adjuvant three times with one-week interval. On the fourth day after the final injection, spleens were extracted from the mice and dispersed to obtain splenocytes.

The splenocytes and SP2/0-Ag14 cells, ATCC CRL-1581, derived from mouse, were co-suspended in a serum-free RPMI 1640 medium, which had been warmed prior to use to 37°C, to give respective cell densities of  $3 \times 10^4$  and  $1 \times 10^4$  cells/ml, and then centrifuged to collect a precipitate. To the precipitate, one milliliter of a serum-free RPMI 1640 medium (pH 7.2) containing 50%(w/v) polyethylene glycol with an average molecular weight of about 15,000 daltons was dropped over one minute, and the resulting mixture was incubated at 37°C for one minute. A serum-free RPMI 1640 medium (pH 7.2) was further dropped to the mixture to give a final volume of 50 ml, which was then centrifuged to collect a precipitate. The precipitate was suspended in HAT

medium, distributed to wells of 96-well microplates in a volume of 200  $\mu$ l/well, and incubated at 37°C for a week to select hybridomas.

Antibodies secreted in culture supernatants in the wells were tested for a reactivity with a Sonic hedgehog protein, obtained in Example 4-1(b), by conventional enzyme-immunoassay to select hybridomas which exhibited the reactivity. The antibodies secreted in the culture supernatants of the selected hybridomas were further tested for another reactivity with the hedgehog protein of this invention, obtained in Example 3, by conventional enzyme-immunoassay to select hybridomas which additionally exhibited the reactivity. Thereafter, the finally selected hybridomas were repeatedly subjected to limiting dilution method, resulting in obtaining hybridoma clones capable of producing the monoclonal antibody of this invention which were named "SH2-3", "SH2-21", and "SH2-260".

#### Example 4-3

##### Production of monoclonal antibody

Hybridomas "SH2-3", "SH2-21", and "SH2-260" obtained in Example 4-2 were separately suspended to give a cell density of  $1 \times 10^5$  cells/ml each in aliquotes of RPMI 1640 medium (pH 7.2) supplemented with 5%(v/v) fetal bovine serum, and cultured at 37°C in a 5%(v/v) CO<sub>2</sub> incubator while scaling up the culture. After the cell densities reached a desired level, the hybridomas were peritoneally injected in a dose of  $1 \times 10^7$  cells/body to eight-week-old BALB/c mice which had been peritoneally injected with 0.5 ml/body "PRISTANE", a reagent of 2,6,10,14-tetramethylpentadecane commercialized by Aldrich Chemical Co., Inc., Milwaukee, USA, and the mice were fed for a week in usual

manner.

From the respective lines of mice, ascites were collected and threefold diluted with PBS. To the dilutions, ammonium sulfate was added to give 50% saturation. The resulting mixture was allowed to stand at 4°C for 24 hours and then centrifuged to collect precipitates. The precipitates were dialyzed against 20 mM KH<sub>2</sub>PO<sub>4</sub> (pH 6.7) at 4°C overnight, and then charged to columns of hydroxyapatite, pre-equilibrated with 20 mM KH<sub>2</sub>PO<sub>4</sub> (pH 6.7). Through the columns, running KH<sub>2</sub>PO<sub>4</sub> (pH 6.7) solution with increasing concentration from 20 to 300 mM in a linear gradient manner resulted in obtaining aqueous solutions of "SH2-3mAb", "SH2-21mAb", and "SH2-260mAb", the monoclonal antibodies of this invention. The yields were about 5 mg/mouse each. Analyzing in usual manner, all of the monoclonal antibodies belonged to a class of IgG<sub>1</sub>.

#### Example 5

##### Western blotting

One microgram of a Desert hedgehog protein, obtained by the method in Example 3, was subjected to SDS-PAGE with 15%(w/v) gel in the presence of a reducing agent. In parallel, 50 ng of a Sonic hedgehog protein, obtained by the method in Example 4-1(b), was subjected to SDS-PAGE with 13%(w/v) gel in the presence of a reducing agent. In usual manner, proteinaceous components in the gels were transferred to nitrocellulose membranes, which were then immersed in "BLOCK ACE™", an immobilizing agent commercialized by Dainippon Pharmaceutical Co., Ltd., Osaka, Japan, to effect blocking. The membranes were immersed in PBS containing 20 µg/ml "SH2-3mAb", monoclonal antibody obtained by the method in Example 4-3, 10%(v/v) "BLOCK



ACE™, and 0.1%(v/v) "TWEEN20", a detergent commercialized by City Chemical Corp., New York, U.S.A., for one hour; and washed with PBS containing 0.1%(v/v) "TWEEN 20" to remove excessive antibodies. Thereafter, the nitrocellulose membranes were reacted for one hour in PBS containing 0.1%(v/v) sheep anti-mouse immunoglobulin antibody labelled with horseradish peroxidase, 10%(v/v) "BLOCK ACE™", and 0.05%(v/v) "TWEEN 20"; washed with PBS containing 0.1%(v/v) "TWEEN 20"; and color-developed by using "ECL™ KIT", a kit for color development commercialized by Amersham International plc, Buckinghamshire, UK. The molecular weight markers used were "SDS-PAGE STANDARDS, LOW RANGE", containing six proteins having distinctive molecular weights of 14,400-97,400 daltons, commercialized by Bio-rad Laboratories Inc., Richmond, USA. The results are in FIG.2.

In FIG.2, on Lane 1, the band corresponding to a molecular weight of  $22,000 \pm 2,000$  is of the hedgehog protein of this invention, and the other band, corresponding to a molecular weight of  $18,000 \pm 2,000$ , is of the degradation product of the hedgehog protein formed during the process in Example 3. In FIG.2, on Lane 2, the band corresponding to a molecular weight of  $22,000 \pm 2,000$  is of a human Sonic hedgehog protein, obtained by the method in Example 4-1(b).

Another Western blotting which was conducted in the same manner as above except for using a monoclonal antibody "SH2-21mAb", obtained by the method in Example 4-3, in place of the monoclonal antibody "SH2-3mAb", giving similar results as above. These results indicate that the monoclonal antibodies, according to this invention, well recognized not only a human Sonic hedgehog protein but also a human Desert hedgehog protein.

## Example 6

### Enzyme-immunoassay

Monoclonal antibodies "SH2-3mAb" and "SH2-260mAb", obtained by the method in Example 4-3, were co-diluted in PBS to give a concentration of 10 µg/ml each, the resulting solution was distributed to wells of 96-well microplates in a volume of 100 µl/well. The microplates were incubated at ambient temperature. From the microplates the solution was removed, and PBS containing 1%(w/v) bovine serum albumin was distributed to the wells in a volume of 200 µl/well. Then the microplates were allowed to stand at 4°C overnight. In parallel, a human Desert hedgehog protein, obtained by the method in Example 3, and a human Sonic hedgehog protein, obtained by the method in Example 4-1(b), were separately diluted with PBS to give desired different concentrations. After removing the solution from the microplates, and the respective hedgehog protein solutions were added to the wells and reacted at ambient temperature for one hour. The microplates were washed with PBS containing 0.05%(v/v) "TWEEN 20", and added with a rabbit anti-hedgehog protein antiserum 500-fold diluted with PBS in a volume of 100 µl/well. The antiserum used in this Example was obtained by immunizing rabbits with a human Sonic hedgehog protein, obtained by the method in Example 4-1(b), and collecting serum from the rabbits in usual manner.

After the reaction with the antiserum, the microplates were washed with PBS containing 0.05%(v/v) "TWEEN 20" and added with a horseradish peroxidase-labelled donkey anti-rabbit immunoglobulin antibody, commercialized by Amersham International plc, Buckinghamshire, UK, which had been 1000-fold diluted with

PBS, followed by allowing the microplates to stand at ambient temperature for one hour. The microplates were washed with PBS containing 0.05%(v/v) "TWEEN 20". Thereafter, in usual manner, a mixture solution of o-phenylene diamine as a substrate and  $H_2O_2$  was added to the wells in a volume of 100  $\mu$ l/well followed by an incubation at ambient temperature for 15 minutes to effect enzyme reaction, and the reaction was terminated by 2N  $H_2SO_4$  added. Intensities of colors in the wells developed by the reaction were estimated by measuring the absorbance at 492 nm. The results are in FIG.3.

The results in FIG.3. indicate that the method for detecting, according to this invention, well detected not only a human Sonic hedgehog protein but also a human Desert hedgehog protein.

As described above, this invention was established based on the finding of a novel hedgehog protein, i.e., a Desert hedgehog protein of human origin. The hedgehog protein of this invention is useful in establishment of a hybridoma capable of producing a monoclonal antibody that recognizes the protein. The hedgehog protein of this invention has efficacy in treatment and prevention of susceptible diseases to the hedgehog protein. The monoclonal antibody is useful in purification and detection of human Desert hedgehog protein because the antibody recognizes the hedgehog protein. The monoclonal antibody has efficacy in treatment, prevention, and diagnosis of diseases relating to excessive production of the hedgehog protein in living bodies. In addition to these effectiveness, the protein, DNA, and monoclonal antibody of this invention are extremely useful in

elucidation of the process of exhibiting hereditary morphological abnormalities in humans. The process of this invention does satisfactorily produce the hedgehog protein.

This invention, which exhibits these remarkable effects, would be very significant and contributive to the art.

While there has been described what is at present considered to be the preferred embodiments of the present invention, it will be understood the various modifications may be made therein, and it is intended to cover in the appended claims all such modifications as fall within the true spirits and scope of the invention.

## SEQUENCE LISTING

### (1) GENERAL INFORMATION:

- (i) APPLICANT: KABUSHIKI KAISHA HAYASHIBARA SEIBUTSU KAGAKU  
KENKYUJO
- (ii) TITLE OF INVENTION: HEDGEHOG PROTEIN
- (iii) NUMBER OF SEQUENCES: 31
- (iv) CORRESPONDENCE ADDRESS:
  - (A) ADDRESSEE: BROWDY AND NEIMARK
  - (B) STREET: 419 Seventh Street N.W., Ste. 300
  - (C) CITY: Washington
  - (D) STATE: D.C.
  - (E) COUNTRY: United States of America
  - (F) ZIP: 20004
- (v) COMPUTER READABLE FORM:
  - (A) MEDIUM TYPE: Floppy disk
  - (B) COMPUTER: IBM PC compatible
  - (C) OPERATING SYSTEM: PC-DOS/MS-DOS
  - (D) SOFTWARE: PatentIn Release #1.0, Version #1.30
- (vi) CURRENT APPLICATION DATA:
  - (A) APPLICATION NUMBER:
  - (B) FILING DATE:
  - (C) CLASSIFICATION:
- (vii) PRIOR APPLICATION DATA:
  - (A) APPLICATION NUMBER: JP 97-121578
  - (B) FILING DATE: 25-APRIL-1997
- (vii) PRIOR APPLICATION DATA:
  - (A) APPLICATION NUMBER: JP 98-
  - (B) FILING DATE: 14-APRIL-1998
- (viii) ATTORNEY/AGENT INFORMATION:
  - (A) NAME: Browdy, Roger L.
  - (B) REGISTRATION NUMBER: 25,618
  - (C) REFERENCE/DOCKET NUMBER:
- (ix) TELECOMMUNICATION INFORMATION:
  - (A) TELEPHONE: (202) 628-5197
  - (B) TELEFAX: (202) 737-35281

### (2) INFORMATION FOR SEQ ID NO: 1:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 176 amino acids
  - (B) TYPE: amino acid
  - (C) strandedness: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

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Gln	Leu	Val	Pro	Leu	Leu	Tyr	Lys	Gln	Phe	Val	Pro	Gly	Val	Pro	Glu
			20					25					30		
Arg	Thr	Leu	Gly	Ala	Ser	Gly	Pro	Ala	Glu	Gly	Arg	Val	Ala	Arg	Gly
		35					40					45			
Ser	Glu	Arg	Phe	Arg	Asp	Leu	Val	Pro	Asn	Tyr	Asn	Pro	Asp	Ile	Ile
	50					55					60				
Phe	Lys	Asp	Glu	Glu	Asn	Ser	Gly	Ala	Asp	Arg	Leu	Met	Thr	Glu	Arg
65					70					75					80
Cys	Lys	Glu	Arg	Val	Asn	Ala	Leu	Ala	Ile	Ala	Val	Met	Asn	Met	Trp
				85					90					95	
Pro	Gly	Val	Arg	Leu	Arg	Val	Thr	Glu	Gly	Trp	Asp	Glu	Asp	Gly	His
			100					105					110		
His	Ala	Gln	Asp	Ser	Leu	His	Tyr	Glu	Gly	Arg	Ala	Leu	Asp	Ile	Thr
		115					120					125			
Thr	Ser	Asp	Arg	Asp	Arg	Asn	Lys	Tyr	Gly	Leu	Leu	Ala	Arg	Leu	Ala
	130					135					140				
Val	Glu	Ala	Gly	Phe	Asp	Trp	Val	Tyr	Tyr	Glu	Ser	Arg	Asn	His	Ile
145					150					155					160
His	Val	Ser	Val	Lys	Ala	Asp	Asn	Ser	Leu	Ala	Val	Arg	Ala	Gly	Gly
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(2) INFORMATION FOR SEQ ID NO: 2:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 374 amino acids
- (B) TYPE: amino acid
- (C) strandedness: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(ix) FEATURE:

- (A) NAME/KEY: mat peptide
- (B) LOCATION: 1..176
- (C) IDENTIFICATION METHOD: S

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:

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Gln	Leu	Val	Pro	Leu	Leu	Tyr	Lys	Gln	Phe	Val	Pro	Gly	Val	Pro	Glu
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Arg	Thr	Leu	Gly	Ala	Ser	Gly	Pro	Ala	Glu	Gly	Arg	Val	Ala	Arg	Gly	
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Ser	Glu	Arg	Phe	Arg	Asp	Leu	Val	Pro	Asn	Tyr	Asn	Pro	Asp	Ile	Ile	
	50					55					60					
Phe	Lys	Asp	Glu	Glu	Asn	Ser	Gly	Ala	Asp	Arg	Leu	Met	Thr	Glu	Arg	
	65				70					75					80	
Cys	Lys	Glu	Arg	Val	Asn	Ala	Leu	Ala	Ile	Ala	Val	Met	Asn	Met	Trp	
				85					90					95		
Pro	Gly	Val	Arg	Leu	Arg	Val	Thr	Glu	Gly	Trp	Asp	Glu	Asp	Gly	His	
			100					105					110			
His	Ala	Gln	Asp	Ser	Leu	His	Tyr	Glu	Gly	Arg	Ala	Leu	Asp	Ile	Thr	
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Thr	Ser	Asp	Arg	Asp	Arg	Asn	Lys	Tyr	Gly	Leu	Leu	Ala	Arg	Leu	Ala	
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Val	Glu	Ala	Gly	Phe	Asp	Trp	Val	Tyr	Tyr	Glu	Ser	Arg	Asn	His	Ile	
	145				150					155					160	
His	Val	Ser	Val	Lys	Ala	Asp	Asn	Ser	Leu	Ala	Val	Arg	Ala	Gly	Gly	
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Cys	Phe	Pro	Gly	Asn	Ala	Thr	Val	Arg	Leu	Trp	Ser	Gly	Glu	Arg	Lys	
			180					185					190			
Gly	Leu	Arg	Glu	Leu	His	Arg	Gly	Asp	Trp	Val	Leu	Thr	Ala	Asp	Ala	
		195					200					205				
Ser	Gly	Arg	Val	Val	Pro	Thr	Pro	Val	Leu	Leu	Phe	Leu	Asp	Arg	Asp	
	210					215					220					
Leu	Gln	Arg	Arg	Ala	Ser	Phe	Val	Ala	Val	Glu	Thr	Glu	Trp	Pro	Pro	
	225				230					235					240	
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Pro	Ala	Pro	Ala	Pro	Gly	Asp	Phe	Ala	Pro	Val	Phe	Ala	Arg	Arg	Leu	
			260					265					270			
Arg	Ala	Gly	Asp	Ser	Val	Leu	Ala	Pro	Gly	Gly	Asp	Ala	Leu	Arg	Pro	
		275					280					285				
Ala	Arg	Val	Ala	Arg	Val	Ala	Arg	Glu	Glu	Ala	Val	Gly	Val	Phe	Ala	
	290					295					300					
Pro	Leu	Thr	Ala	His	Gly	Thr	Leu	Leu	Val	Asn	Asp	Val	Leu	Ala	Ser	
	305				310					315					320	
Cys	Tyr	Ala	Val	Leu	Glu	Ser	His	Gln	Trp	Ala	His	Arg	Ala	Phe	Ala	
				325					330					335		

Pro Leu Arg Leu Leu His Ala Leu Gly Ala Leu Leu Pro Gly Gly Ala  
340 345 350

Val Gln Pro Thr Gly Met His Trp Tyr Ser Arg Leu Leu Tyr Arg Leu  
355 360 365

Ala Glu Glu Leu Leu Gly  
370

(2) INFORMATION FOR SEQ ID NO: 3:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 396 amino acids  
 (B) TYPE: amino acid  
 (C) strandedness: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

- (ix) FEATURE:  
 (A) NAME/KEY: sig peptide  
 (B) LOCATION: -22..-1  
 (C) IDENTIFICATION METHOD: S  
 (A) NAME/KEY: mat peptide  
 (B) LOCATION: 1..176  
 (C) IDENTIFICATION METHOD: S

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3:

Met Ala Leu Leu Thr Asn Leu Leu Pro Leu Cys Cys Leu Ala Leu Leu  
-20 -15 -10

Ala Leu Pro Ala Gln Ser Cys Gly Pro Gly Arg Gly Pro Val Gly Arg  
-5 1 5 10

Arg Arg Tyr Ala Arg Lys Gln Leu Val Pro Leu Leu Tyr Lys Gln Phe  
15 20 25

Val Pro Gly Val Pro Glu Arg Thr Leu Gly Ala Ser Gly Pro Ala Glu  
30 35 40

Gly Arg Val Ala Arg Gly Ser Glu Arg Phe Arg Asp Leu Val Pro Asn  
45 50 55

Tyr Asn Pro Asp Ile Ile Phe Lys Asp Glu Glu Asn Ser Gly Ala Asp  
60 65 70

Arg Leu Met Thr Glu Arg Cys Lys Glu Arg Val Asn Ala Leu Ala Ile  
75 80 85 90

Ala Val Met Asn Met Trp Pro Gly Val Arg Leu Arg Val Thr Glu Gly  
95 100 105

Trp Asp Glu Asp Gly His His Ala Gln Asp Ser Leu His Tyr Glu Gly  
110 115 120

Arg Ala Leu Asp Ile Thr Thr Ser Asp Arg Asp Arg Asn Lys Tyr Gly



125					130					135					
Leu	Leu	Ala	Arg	Leu	Ala	Val	Glu	Ala	Gly	Phe	Asp	Trp	Val	Tyr	Tyr
140						145					150				
Glu	Ser	Arg	Asn	His	Ile	His	Val	Ser	Val	Lys	Ala	Asp	Asn	Ser	Leu
155					160					165					170
Ala	Val	Arg	Ala	Gly	Gly	Cys	Phe	Pro	Gly	Asn	Ala	Thr	Val	Arg	Leu
				175					180					185	
Trp	Ser	Gly	Glu	Arg	Lys	Gly	Leu	Arg	Glu	Leu	His	Arg	Gly	Asp	Trp
			190					195					200		
Val	Leu	Thr	Ala	Asp	Ala	Ser	Gly	Arg	Val	Val	Pro	Thr	Pro	Val	Leu
		205					210					215			
Leu	Phe	Leu	Asp	Arg	Asp	Leu	Gln	Arg	Arg	Ala	Ser	Phe	Val	Ala	Val
	220					225					230				
Glu	Thr	Glu	Trp	Pro	Pro	Arg	Lys	Leu	Leu	Leu	Thr	Pro	Trp	His	Leu
235					240					245					250
Val	Phe	Ala	Ala	Arg	Gly	Pro	Ala	Pro	Ala	Pro	Gly	Asp	Phe	Ala	Pro
				255					260					265	
Val	Phe	Ala	Arg	Arg	Leu	Arg	Ala	Gly	Asp	Ser	Val	Leu	Ala	Pro	Gly
			270					275					280		
Gly	Asp	Ala	Leu	Arg	Pro	Ala	Arg	Val	Ala	Arg	Val	Ala	Arg	Glu	Glu
		285					290					295			
Ala	Val	Gly	Val	Phe	Ala	Pro	Leu	Thr	Ala	His	Gly	Thr	Leu	Leu	Val
		300				305					310				
Asn	Asp	Val	Leu	Ala	Ser	Cys	Tyr	Ala	Val	Leu	Glu	Ser	His	Gln	Trp
315					320					325					330
Ala	His	Arg	Ala	Phe	Ala	Pro	Leu	Arg	Leu	Leu	His	Ala	Leu	Gly	Ala
				335					340					345	
Leu	Leu	Pro	Gly	Gly	Ala	Val	Gln	Pro	Thr	Gly	Met	His	Trp	Tyr	Ser
			350					355					360		
Arg	Leu	Leu	Tyr	Arg	Leu	Ala	Glu	Glu	Leu	Leu	Gly				
		365					370								

(2) INFORMATION FOR SEQ ID NO: 4:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 528 base pairs
  - (B) TYPE: nucleic acid
  - (C) strandedness: double
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(ix) FEATURE:  
 (A) NAME/KEY: mat peptide  
 (B) LOCATION: 1..528  
 (C) IDENTIFICATION METHOD: S

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:

TGC	GGG	CCG	GGC	CGG	GGG	CCG	GTT	GGC	CGG	CGC	CGC	TAT	GCG	CGC	AAG	48
Cys	Gly	Pro	Gly	Arg	Gly	Pro	Val	Gly	Arg	Arg	Arg	Tyr	Ala	Arg	Lys	
1				5					10					15		
CAG	CTC	GTG	CCG	CTA	CTC	TAC	AAG	CAA	TTT	GTG	CCC	GGC	GTG	CCA	GAG	96
Gln	Leu	Val	Pro	Leu	Leu	Tyr	Lys	Gln	Phe	Val	Pro	Gly	Val	Pro	Glu	
			20					25					30			
CGG	ACC	CTG	GGC	GCC	AGT	GGG	CCA	GCG	GAG	GGG	AGG	GTG	GCA	AGG	GGC	144
Arg	Thr	Leu	Gly	Ala	Ser	Gly	Pro	Ala	Glu	Gly	Arg	Val	Ala	Arg	Gly	
		35					40					45				
TCC	GAG	CGC	TTC	CGG	GAC	CTC	GTG	CCC	AAC	TAC	AAC	CCC	GAC	ATC	ATC	192
Ser	Glu	Arg	Phe	Arg	Asp	Leu	Val	Pro	Asn	Tyr	Asn	Pro	Asp	Ile	Ile	
	50					55					60					
TTC	AAG	GAT	GAG	GAG	AAC	AGT	GGA	GCC	GAC	CGC	CTG	ATG	ACC	GAA	CGT	240
Phe	Lys	Asp	Glu	Glu	Asn	Ser	Gly	Ala	Asp	Arg	Leu	Met	Thr	Glu	Arg	
65					70					75				80		
TGT	AAG	GAA	CGG	GTG	AAC	GCT	TTG	GCC	ATT	GCC	GTG	ATG	AAC	ATG	TGG	288
Cys	Lys	Glu	Arg	Val	Asn	Ala	Leu	Ala	Ile	Ala	Val	Met	Asn	Met	Trp	
				85					90					95		
CCC	GGA	GTG	CGC	CTA	CGA	GTG	ACT	GAG	GGC	TGG	GAC	GAG	GAC	GGC	CAC	336
Pro	Gly	Val	Arg	Leu	Arg	Val	Thr	Glu	Gly	Trp	Asp	Glu	Asp	Gly	His	
			100					105					110			
CAC	GCT	CAG	GAT	TCA	CTC	CAC	TAC	GAA	GGC	CGT	GCT	TTG	GAC	ATC	ACT	384
His	Ala	Gln	Asp	Ser	Leu	His	Tyr	Glu	Gly	Arg	Ala	Leu	Asp	Ile	Thr	
		115					120					125				
ACG	TCT	GAC	CGC	GAC	CGC	AAC	AAG	TAT	GGG	TTG	CTG	GCG	CGC	CTC	GCA	432
Thr	Ser	Asp	Arg	Asp	Arg	Asn	Lys	Tyr	Gly	Leu	Leu	Ala	Arg	Leu	Ala	
	130					135					140					
GTG	GAA	GCC	GGC	TTC	GAC	TGG	GTC	TAC	TAC	GAG	TCC	CGC	AAC	CAC	ATC	480
Val	Glu	Ala	Gly	Phe	Asp	Trp	Val	Tyr	Tyr	Glu	Ser	Arg	Asn	His	Ile	
145					150					155					160	
CAC	GTG	TCG	GTC	AAA	GCT	GAT	AAC	TCA	CTG	GCG	GTC	CGG	GCG	GGC	GGC	528
His	Val	Ser	Val	Lys	Ala	Asp	Asn	Ser	Leu	Ala	Val	Arg	Ala	Gly	Gly	
				165					170					175		

(2) INFORMATION FOR SEQ ID NO: 5:

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 1122 base pairs  
 (B) TYPE: nucleic acid  
 (C) strandedness: double

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(ix) FEATURE:

(A) NAME/KEY: mat peptide

(B) LOCATION: 1..528

(C) IDENTIFICATION METHOD: S

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:

TGC	GGG	CCG	GGC	CGG	GGG	CCG	GTT	GGC	CGG	CGC	CGC	TAT	GCG	CGC	AAG	48
Cys	Gly	Pro	Gly	Arg	Gly	Pro	Val	Gly	Arg	Arg	Arg	Tyr	Ala	Arg	Lys	
1				5				10						15		
CAG	CTC	GTG	CCG	CTA	CTC	TAC	AAG	CAA	TTT	GTG	CCC	GGC	GTG	CCA	GAG	96
Gln	Leu	Val	Pro	Leu	Leu	Tyr	Lys	Gln	Phe	Val	Pro	Gly	Val	Pro	Glu	
			20					25					30			
CGG	ACC	CTG	GGC	GCC	AGT	GGG	CCA	GCG	GAG	GGG	AGG	GTG	GCA	AGG	GGC	144
Arg	Thr	Leu	Gly	Ala	Ser	Gly	Pro	Ala	Glu	Gly	Arg	Val	Ala	Arg	Gly	
		35					40					45				
TCC	GAG	CGC	TTC	CGG	GAC	CTC	GTG	CCC	AAC	TAC	AAC	CCC	GAC	ATC	ATC	192
Ser	Glu	Arg	Phe	Arg	Asp	Leu	Val	Pro	Asn	Tyr	Asn	Pro	Asp	Ile	Ile	
	50					55					60					
TTC	AAG	GAT	GAG	GAG	AAC	AGT	GGA	GCC	GAC	CGC	CTG	ATG	ACC	GAA	CGT	240
Phe	Lys	Asp	Glu	Glu	Asn	Ser	Gly	Ala	Asp	Arg	Leu	Met	Thr	Glu	Arg	
	65				70					75					80	
TGT	AAG	GAA	CGG	GTG	AAC	GCT	TTG	GCC	ATT	GCC	GTG	ATG	AAC	ATG	TGG	288
Cys	Lys	Glu	Arg	Val	Asn	Ala	Leu	Ala	Ile	Ala	Val	Met	Asn	Met	Trp	
				85					90					95		
CCC	GGA	GTG	CGC	CTA	CGA	GTG	ACT	GAG	GGC	TGG	GAC	GAG	GAC	GGC	CAC	336
Pro	Gly	Val	Arg	Leu	Arg	Val	Thr	Glu	Gly	Trp	Asp	Glu	Asp	Gly	His	
			100					105					110			
CAC	GCT	CAG	GAT	TCA	CTC	CAC	TAC	GAA	GGC	CGT	GCT	TTG	GAC	ATC	ACT	384
His	Ala	Gln	Asp	Ser	Leu	His	Tyr	Glu	Gly	Arg	Ala	Leu	Asp	Ile	Thr	
		115					120					125				
ACG	TCT	GAC	CGC	GAC	CGC	AAC	AAG	TAT	GGG	TTG	CTG	GCG	CGC	CTC	GCA	432
Thr	Ser	Asp	Arg	Asp	Arg	Asn	Lys	Tyr	Gly	Leu	Leu	Ala	Arg	Leu	Ala	
		130				135					140					
GTG	GAA	GCC	GGC	TTC	GAC	TGG	GTC	TAC	TAC	GAG	TCC	CGC	AAC	CAC	ATC	480
Val	Glu	Ala	Gly	Phe	Asp	Trp	Val	Tyr	Tyr	Glu	Ser	Arg	Asn	His	Ile	
	145				150					155					160	
CAC	GTG	TCG	GTC	AAA	GCT	GAT	AAC	TCA	CTG	GCG	GTC	CGG	GCG	GGC	GGC	528
His	Val	Ser	Val	Lys	Ala	Asp	Asn	Ser	Leu	Ala	Val	Arg	Ala	Gly	Gly	
				165					170					175		
TGC	TTT	CCG	GGA	AAT	GCA	ACT	GTG	CGC	CTG	TGG	AGC	GGC	GAG	CGG	AAA	576
Cys	Phe	Pro	Gly	Asn	Ala	Thr	Val	Arg	Leu	Trp	Ser	Gly	Glu	Arg	Lys	

180						185						190						
GGG	CTG	CGG	GAA	CTG	CAC	CGC	GGA	GAC	TGG	GTT	TTG	ACG	GCC	GAT	GCG	624		
Gly	Leu	Arg	Glu	Leu	His	Arg	Gly	Asp	Trp	Val	Leu	Thr	Ala	Asp	Ala			
		195					200					205						
TCA	GGC	CGG	GTG	GTG	CCC	ACG	CCG	GTG	CTG	CTC	TTC	CTG	GAC	CGG	GAC	672		
Ser	Gly	Arg	Val	Val	Pro	Thr	Pro	Val	Leu	Leu	Phe	Leu	Asp	Arg	Asp			
	210					215					220							
TTG	CAG	CGC	CGG	GCT	TCA	TTT	GTG	GCT	GTG	GAG	ACC	GAG	TGG	CCT	CCA	720		
Leu	Gln	Arg	Arg	Ala	Ser	Phe	Val	Ala	Val	Glu	Thr	Glu	Trp	Pro	Pro			
225					230					235					240			
CGC	AAA	CTG	TTG	CTC	ACG	CCC	TGG	CAC	CTG	GTG	TTT	GCC	GCT	CGA	GGG	768		
Arg	Lys	Leu	Leu	Leu	Thr	Pro	Trp	His	Leu	Val	Phe	Ala	Ala	Arg	Gly			
				245					250					255				
CCG	GCG	CCC	GCG	CCA	GGC	GAC	TTT	GCA	CCG	GTG	TTC	GCG	CGC	CGG	CTA	816		
Pro	Ala	Pro	Ala	Pro	Gly	Asp	Phe	Ala	Pro	Val	Phe	Ala	Arg	Arg	Leu			
			260					265					270					
CGC	GCT	GGG	GAC	TCG	GTG	CTG	GCG	CCC	GGC	GGG	GAT	GCG	CTT	CGG	CCA	864		
Arg	Ala	Gly	Asp	Ser	Val	Leu	Ala	Pro	Gly	Gly	Asp	Ala	Leu	Arg	Pro			
		275					280					285						
GCG	CGC	GTG	GCC	CGT	GTG	GCG	CGG	GAG	GAA	GCC	GTG	GCG	GTG	TTC	GCG	912		
Ala	Arg	Val	Ala	Arg	Val	Ala	Arg	Glu	Glu	Ala	Val	Gly	Val	Phe	Ala			
	290					295					300							
CCG	CTC	ACC	GCG	CAC	GGG	ACG	CTG	CTG	GTG	AAC	GAT	GTC	CTG	GCC	TCT	960		
Pro	Leu	Thr	Ala	His	Gly	Thr	Leu	Leu	Val	Asn	Asp	Val	Leu	Ala	Ser			
					310					315					320			
TGC	TAC	GCG	GTT	CTG	GAG	AGT	CAC	CAG	TGG	GCG	CAC	CGC	GCT	TTT	GCC	1008		
Cys	Tyr	Ala	Val	Leu	Glu	Ser	His	Gln	Trp	Ala	His	Arg	Ala	Phe	Ala			
				325					330					335				
CCC	TTG	AGA	CTG	CTG	CAC	GCG	CTA	GGG	GCG	CTG	CTC	CCC	GGC	GGG	GCC	1056		
Pro	Leu	Arg	Leu	Leu	His	Ala	Leu	Gly	Ala	Leu	Leu	Pro	Gly	Gly	Ala			
			340					345					350					
GTC	CAG	CCG	ACT	GGC	ATG	CAT	TGG	TAC	TCT	CGG	CTC	CTC	TAC	CGC	TTA	1104		
Val	Gln	Pro	Thr	Gly	Met	His	Trp	Tyr	Ser	Arg	Leu	Leu	Tyr	Arg	Leu			
		355					360					365						
GCG	GAG	GAG	CTA	CTG	GGC											1122		
Ala	Glu	Glu	Leu	Leu	Gly													
					370													

(2) INFORMATION FOR SEQ ID NO: 6:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1188 base pairs
- (B) TYPE: nucleic acid
- (C) strandedness: double
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(ix) FEATURE:

(A) NAME/KEY: sig peptide  
(B) LOCATION: 1..66  
(C) IDENTIFICATION METHOD: S  
(A) NAME/KEY: mat peptide  
(B) LOCATION: 67..594  
(C) IDENTIFICATION METHOD: S

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:

ATG	GCT	CTC	CTG	ACC	AAT	CTA	CTG	CCC	CTG	TGC	TGC	TTG	GCA	CTT	CTG	48
Met	Ala	Leu	Leu	Thr	Asn	Leu	Leu	Pro	Leu	Cys	Cys	Leu	Ala	Leu	Leu	
		-20					-15					-10				
GCG	CTG	CCA	GCC	CAG	AGC	TGC	GGG	CCG	GGC	CGG	GGG	CCG	GTT	GGC	CGG	96
Ala	Leu	Pro	Ala	Gln	Ser	Cys	Gly	Pro	Gly	Arg	Gly	Pro	Val	Gly	Arg	
	-5					1				5					10	
CGC	CGC	TAT	GCG	CGC	AAG	CAG	CTC	GTG	CCG	CTA	CTC	TAC	AAG	CAA	TTT	144
Arg	Arg	Tyr	Ala	Arg	Lys	Gln	Leu	Val	Pro	Leu	Leu	Tyr	Lys	Gln	Phe	
				15					20					25		
GTG	CCC	GGC	GTG	CCA	GAG	CGG	ACC	CTG	GGC	GCC	AGT	GGG	CCA	GCG	GAG	192
Val	Pro	Gly	Val	Pro	Glu	Arg	Thr	Leu	Gly	Ala	Ser	Gly	Pro	Ala	Glu	
			30					35					40			
GGG	AGG	GTG	GCA	AGG	GGC	TCC	GAG	CGC	TTC	CGG	GAC	CTC	GTG	CCC	AAC	240
Gly	Arg	Val	Ala	Arg	Gly	Ser	Glu	Arg	Phe	Arg	Asp	Leu	Val	Pro	Asn	
		45					50					55				
TAC	AAC	CCC	GAC	ATC	ATC	TTC	AAG	GAT	GAG	GAG	AAC	AGT	GGA	GCC	GAC	288
Tyr	Asn	Pro	Asp	Ile	Ile	Phe	Lys	Asp	Glu	Glu	Asn	Ser	Gly	Ala	Asp	
	60					65					70					
CGC	CTG	ATG	ACC	GAA	CGT	TGT	AAG	GAA	CGG	GTG	AAC	GCT	TTG	GCC	ATT	336
Arg	Leu	Met	Thr	Glu	Arg	Cys	Lys	Glu	Arg	Val	Asn	Ala	Leu	Ala	Ile	
	75				80					85					90	
GCC	GTG	ATG	AAC	ATG	TGG	CCC	GGA	GTG	CGC	CTA	CGA	GTG	ACT	GAG	GGC	384
Ala	Val	Met	Asn	Met	Trp	Pro	Gly	Val	Arg	Leu	Arg	Val	Thr	Glu	Gly	
				95				100						105		
TGG	GAC	GAG	GAC	GGC	CAC	CAC	GCT	CAG	GAT	TCA	CTC	CAC	TAC	GAA	GGC	432
Trp	Asp	Glu	Asp	Gly	His	His	Ala	Gln	Asp	Ser	Leu	His	Tyr	Glu	Gly	
			110					115					120			
CGT	GCT	TTG	GAC	ATC	ACT	ACG	TCT	GAC	CGC	GAC	CGC	AAC	AAG	TAT	GGG	480
Arg	Ala	Leu	Asp	Ile	Thr	Thr	Ser	Asp	Arg	Asp	Arg	Asn	Lys	Tyr	Gly	
		125					130					135				
TTG	CTG	GCG	CGC	CTC	GCA	GTG	GAA	GCC	GGC	TTC	GAC	TGG	GTC	TAC	TAC	528
Leu	Leu	Ala	Arg	Leu	Ala	Val	Glu	Ala	Gly	Phe	Asp	Trp	Val	Tyr	Tyr	
	140					145				150						
GAG	TCC	CGC	AAC	CAC	ATC	CAC	GTG	TCG	GTC	AAA	GCT	GAT	AAC	TCA	CTG	576

Glu 155	Ser	Arg	Asn	His	Ile 160	His	Val	Ser	Val	Lys 165	Ala	Asp	Asn	Ser	Leu 170	
GCG Ala	GTC Val	CGG Arg	GCG Ala	GGC Gly 175	GGC Gly	TGC Cys	TTT Phe	CCG Pro	GGA Gly 180	AAT Asn	GCA Ala	ACT Thr	GTG Val	CGC Arg 185	CTG Leu	624
TGG Trp	AGC Ser	GGC Gly	GAG Glu 190	CGG Arg	AAA Lys	GGG Gly	CTG Leu	CGG Arg 195	GAA Glu	CTG Leu	CAC His	CGC Arg	GGA Gly 200	GAC Asp	TGG Trp	672
GTT Val	TTG Leu	ACG Thr 205	GCC Ala	GAT Asp	GCG Ala	TCA Ser	GGC Gly 210	CGG Arg	GTG Val	GTG Val	CCC Pro	ACG Thr 215	CCG Pro	GTG Val	CTG Leu	720
CTC Leu	TTC Phe 220	CTG Leu	GAC Asp	CGG Arg	GAC Asp	TTG Leu 225	CAG Gln	CGC Arg	CGG Arg	GCT Ala	TCA Ser 230	TTT Phe	GTG Val	GCT Ala	GTG Val	768
GAG Glu 235	ACC Thr	GAG Glu	TGG Trp	CCT Pro	CCA Pro 240	CGC Arg	AAA Lys	CTG Leu	TTG Leu	CTC Leu 245	ACG Thr	CCC Pro	TGG Trp	CAC His	CTG Leu 250	816
GTG Val	TTT Phe	GCC Ala	GCT Ala	CGA Arg 255	GGG Gly	CCG Pro	GCG Ala	CCC Pro	GCG Ala 260	CCA Pro	GGC Gly	GAC Asp	TTT Phe	GCA Ala 265	CCG Pro	864
GTG Val	TTC Phe	GCG Ala	CGC Arg 270	CGG Arg	CTA Leu	CGC Arg	GCT Ala	GGG Gly 275	GAC Asp	TCG Ser	GTG Val	CTG Leu	GCG Ala 280	CCC Pro	GGC Gly	912
GGG Gly	GAT Asp	GCG Ala 285	CTT Leu	CGG Arg	CCA Pro	GCG Ala	CGC Arg 290	GTG Val	GCC Ala	CGT Arg	GTG Val	GCG Ala 295	CGG Arg	GAG Glu	GAA Glu	960
GCC Ala 300	GTG Val	GGC Gly	GTG Val	TTC Phe	GCG Ala	CCG Pro 305	CTC Leu	ACC Thr	GCG Ala	CAC His	GGG Gly 310	ACG Thr	CTG Leu	CTG Leu	GTG Val	1008
AAC Asn 315	GAT Asp	GTC Val	CTG Leu	GCC Ala	TCT Ser 320	TGC Cys	TAC Tyr	GCG Ala	GTT Val	CTG Leu 325	GAG Glu	AGT Ser	CAC His	CAG Gln	TGG Trp 330	1056
GCG Ala	CAC His	CGC Arg	GCT Ala	TTT Phe 335	GCC Ala	CCC Pro	TTG Leu	AGA Arg	CTG Leu 340	CTG Leu	CAC His	GCG Ala	CTA Leu	GGG Gly 345	GCG Ala	1104
CTG Leu	CTC Leu	CCC Pro	GGC Gly 350	GGG Gly	GCC Ala	GTC Val	CAG Gln 355	CCG Pro	ACT Thr	GGC Gly	ATG Met	CAT His	TGG Trp 360	TAC Tyr	TCT Ser	1152
CGG Arg	CTC Leu	CTC Leu	TAC Tyr 365	CGC Arg	TTA Leu	GCG Ala	GAG Glu 370	GAG Glu	CTA Leu	CTG Leu	GGC Gly					1188

(2) INFORMATION FOR SEQ ID NO: 7:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 548 base pairs
- (B) TYPE: nucleic acid
- (C) strandedness: double
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: human
- (B) INDIVIDUAL ISOLATE: ARH-77, ATCC CRL-1621

(ix) FEATURE:

- (A) NAME/KEY: sig peptide
- (B) LOCATION: 1..18
- (C) IDENTIFICATION METHOD: S
- (A) NAME/KEY: mat peptide
- (B) LOCATION: 19..546
- (C) IDENTIFICATION METHOD: S

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:

GCG	CTG	CCA	GCC	CAG	AGC	TGC	GGG	CCG	GGC	CGG	GGG	CCG	GTT	GGC	CGG	48
Ala	Leu	Pro	Ala	Gln	Ser	Cys	Gly	Pro	Gly	Arg	Gly	Pro	Val	Gly	Arg	
	-5					1				5					10	
CGC	CGC	TAT	GCG	CGC	AAG	CAG	CTC	GTG	CCG	CTA	CTC	TAC	AAG	CAA	TTT	96
Arg	Arg	Tyr	Ala	Arg	Lys	Gln	Leu	Val	Pro	Leu	Leu	Tyr	Lys	Gln	Phe	
				15					20					25		
GTG	CCC	GGC	GTG	CCA	GAG	CGG	ACC	CTG	GGC	GCC	AGT	GGG	CCA	GCG	GAG	144
Val	Pro	Gly	Val	Pro	Glu	Arg	Thr	Leu	Gly	Ala	Ser	Gly	Pro	Ala	Glu	
			30					35					40			
GGG	AGG	GTG	GCA	AGG	GGC	TCC	GAG	CGC	TTC	CGG	GAC	CTC	GTG	CCC	AAC	192
Gly	Arg	Val	Ala	Arg	Gly	Ser	Glu	Arg	Phe	Arg	Asp	Leu	Val	Pro	Asn	
		45					50					55				
TAC	AAC	CCC	GAC	ATC	ATC	TTC	AAG	GAT	GAG	GAG	AAC	AGT	GGA	GCC	GAC	240
Tyr	Asn	Pro	Asp	Ile	Ile	Phe	Lys	Asp	Glu	Glu	Asn	Ser	Gly	Ala	Asp	
	60					65					70					
CGC	CTG	ATG	ACC	GAA	CGT	TGT	AAG	GAA	CGG	GTG	AAC	GCT	TTG	GCC	ATT	288
Arg	Leu	Met	Thr	Glu	Arg	Cys	Lys	Glu	Arg	Val	Asn	Ala	Leu	Ala	Ile	
	75				80					85					90	
GCC	GTG	ATG	AAC	ATG	TGG	CCC	GGA	GTG	CGC	CTA	CGA	GTG	ACT	GAG	GGC	336
Ala	Val	Met	Asn	Met	Trp	Pro	Gly	Val	Arg	Leu	Arg	Val	Thr	Glu	Gly	
				95				100						105		
TGG	GAC	GAG	GAC	GGC	CAC	CAC	GCT	CAG	GAT	TCA	CTC	CAC	TAC	GAA	GGC	384
Trp	Asp	Glu	Asp	Gly	His	His	Ala	Gln	Asp	Ser	Leu	His	Tyr	Glu	Gly	
			110					115					120			
CGT	GCT	TTG	GAC	ATC	ACT	ACG	TCT	GAC	CGC	GAC	CGC	AAC	AAG	TAT	GGG	432
Arg	Ala	Leu	Asp	Ile	Thr	Thr	Ser	Asp	Arg	Asp	Arg	Asn	Lys	Tyr	Gly	
		125					130					135				

TTG	CTG	GCG	CGC	CTC	GCA	GTG	GAA	GCC	GGC	TTC	GAC	TGG	GTC	TAC	TAC	480
Leu	Leu	Ala	Arg	Leu	Ala	Val	Glu	Ala	Gly	Phe	Asp	Trp	Val	Tyr	Tyr	
	140					145					150					
GAG	TCC	CGC	AAC	CAC	ATC	CAC	GTG	TCG	GTC	AAA	GCT	GAT	AAC	TCA	CTG	528
Glu	Ser	Arg	Asn	His	Ile	His	Val	Ser	Val	Lys	Ala	Asp	Asn	Ser	Leu	
	155				160					165					170	
GCG	GTC	CGG	GCG	GGC	GGC	TG										548
Ala	Val	Arg	Ala	Gly	Gly											
				175												

(2) INFORMATION FOR SEQ ID NO: 8:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 602 base pairs
- (B) TYPE: nucleic acid
- (C) strandedness: double
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: human
- (B) INDIVIDUAL ISOLATE: ARH-77, ATCC CRL-1621

(ix) FEATURE:

- (A) NAME/KEY: 5'UTR
- (B) LOCATION: 1..6
- (C) IDENTIFICATION METHOD: S
- (A) NAME/KEY: sig peptide
- (B) LOCATION: 7..72
- (C) IDENTIFICATION METHOD: S
- (A) NAME/KEY: mat peptide
- (B) LOCATION: 73..600
- (C) IDENTIFICATION METHOD: S

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 8:

GTATCC	ATG	GCT	CTC	CTG	ACC	AAT	CTA	CTG	CCC	CTG	TGC	TGC	TTG	GCA	48	
	Met	Ala	Leu	Leu	Thr	Asn	Leu	Leu	Pro	Leu	Cys	Cys	Leu	Ala		
			-20					-15					-10			
CTT	CTG	GCG	CTG	CCA	GCC	CAG	AGC	TGC	GGG	CCG	GGC	CGG	GGG	CCG	GTT	96
Leu	Leu	Ala	Leu	Pro	Ala	Gln	Ser	Cys	Gly	Pro	Gly	Arg	Gly	Pro	Val	
			-5					1				5				
GGC	CGG	CGC	CGC	TAT	GCG	CGC	AAG	CAG	CTC	GTG	CCG	CTA	CTC	TAC	AAG	144
Gly	Arg	Arg	Arg	Tyr	Ala	Arg	Lys	Gln	Leu	Val	Pro	Leu	Leu	Tyr	Lys	
	10					15					20					
CAA	TTT	GTG	CCC	GGC	GTG	CCA	GAG	CGG	ACC	CTG	GGC	GCC	AGT	GGG	CCA	192
Gln	Phe	Val	Pro	Gly	Val	Pro	Glu	Arg	Thr	Leu	Gly	Ala	Ser	Gly	Pro	
	25				30					35					40	
GCG	GAG	GGG	AGG	GTG	GCA	AGG	GGC	TCC	GAG	CGC	TTC	CGG	GAC	CTC	GTG	240
Ala	Glu	Gly	Arg	Val	Ala	Arg	Gly	Ser	Glu	Arg	Phe	Arg	Asp	Leu	Val	



				45				50				55					
CCC Pro	AAC Asn	TAC Tyr	AAC Asn 60	CCC Pro	GAC Asp	ATC Ile	ATC Ile	TTC Phe 65	AAG Lys	GAT Asp	GAG Glu	GAG Glu	AAC Asn 70	AGT Ser	GGA Gly	288	
GCC Ala	GAC Asp	CGC Arg 75	CTG Leu	ATG Met	ACC Thr	GAA Glu	CGT Arg 80	TGT Cys	AAG Lys	GAA Glu	CGG Arg	GTG Val 85	AAC Asn	GCT Ala	TTG Leu	336	
GCC Ala	ATT Ile 90	GCC Ala	GTG Val	ATG Met	AAC Asn	ATG Met 95	TGG Trp	CCC Pro	GGA Gly	GTG Val	CGC Arg 100	CTA Leu	CGA Arg	GTG Val	ACT Thr	384	
GAG Glu 105	GGC Gly	TGG Trp	GAC Asp	GAG Glu	GAC Asp 110	GGC Gly	CAC His	CAC His	GCT Ala	CAG Gln 115	GAT Asp	TCA Ser	CTC Leu	CAC His	TAC Tyr 120	432	
GAA Glu	GGC Gly	CGT Arg	GCT Ala	TTG Leu 125	GAC Asp	ATC Ile	ACT Thr	ACG Thr	TCT Ser 130	GAC Asp	CGC Arg	GAC Asp	CGC Arg	AAC Asn 135	AAG Lys	480	
TAT Tyr	GGG Gly	TTG Leu	CTG Leu 140	GCG Ala	CGC Arg	CTC Leu	GCA Ala	GTG Val 145	GAA Glu	GCC Ala	GGC Gly	TTC Phe	GAC Asp 150	TGG Trp	GTC Val	528	
TAC Tyr	TAC Tyr	GAG Glu 155	TCC Ser	CGC Arg	AAC Asn	CAC His	ATC Ile 160	CAC His	GTG Val	TCG Ser	GTC Val	AAA Lys 165	GCT Ala	GAT Asp	AAC Asn	576	
TCA Ser	CTG Leu 170	GCG Ala	GTC Val	CGG Arg	GCG Ala	GGC Gly 175	GGC Gly	TG								602	

(2) INFORMATION FOR SEQ ID NO: 9:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 575 base pairs  
(B) TYPE: nucleic acid  
(C) strandedness: double  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: human  
(B) INDIVIDUAL ISOLATE: ARH-77, ATCC CRL-1621

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9:

C	GTG	TCG	GTC	AAA	GCT	GAT	AAC	TCA	CTG	GCG	GTC	CGG	GCG	GGC	GGC	46
	Val	Ser	Val	Lys	Ala	Asp	Asn	Ser	Leu	Ala	Val	Arg	Ala	Gly	Gly	
	1				5					10					15	
TGC	TTT	CCG	GGA	AAT	GCA	ACT	GTG	CGC	CTG	TGG	AGC	GGC	GAG	CGG	AAA	94
Cys	Phe	Pro	Gly	Asn	Ala	Thr	Val	Arg	Leu	Trp	Ser	Gly	Glu	Arg	Lys	
				20					25						30	

[illegible]

(2) INFORMATION FOR SEQ ID NO: 10:

- (A) LENGTH: 230 base pairs  
(B) TYPE: nucleic acid  
(C) strandedness: double  
(D) TOPOLOGY: linear

(vi) ORIGINAL SOURCE:

- (B) INDIVIDUAL ISOLATE: ARH-77, ATCC CRL-1621

(A) NAME/KEY: 3'UTR  
 (B) LOCATION: 218..230  
 (C) IDENTIFICATION METHOD: S

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 10:

G	TTC	GCG	CCG	CTC	ACC	GCG	CAC	GGG	ACG	CTG	CTG	GTG	AAC	GAT	GTC	46
Phe	Ala	Pro	Leu	Thr	Ala	His	Gly	Thr	Leu	Leu	Val	Asn	Asp	Val		
1					5				10					15		
CTG	GCC	TCT	TGC	TAC	GCG	GTT	CTG	GAG	AGT	CAC	CAG	TGG	GCG	CAC	CGC	94
Leu	Ala	Ser	Cys	Tyr	Ala	Val	Leu	Glu	Ser	His	Gln	Trp	Ala	His	Arg	
				20					25					30		
GCT	TTT	GCC	CCC	TTG	AGA	CTG	CTG	CAC	GCG	CTA	GGG	GCG	CTG	CTC	CCC	142
Ala	Phe	Ala	Pro	Leu	Arg	Leu	Leu	His	Ala	Leu	Gly	Ala	Leu	Leu	Pro	
			35					40					45			
GGC	GGG	GCC	GTC	CAG	CCG	ACT	GGC	ATG	CAT	TGG	TAC	TCT	CGG	CTC	CTC	190
Gly	Gly	Ala	Val	Gln	Pro	Thr	Gly	Met	His	Trp	Tyr	Ser	Arg	Leu	Leu	
		50					55					60				
TAC	CGC	TTA	GCG	GAG	GAG	CTA	CTG	GGC	TGAGCGTCCC	AGG						230
Tyr	Arg	Leu	Ala	Glu	Glu	Leu	Leu	Gly								
	65					70										

(2) INFORMATION FOR SEQ ID NO: 11:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 522 base pairs  
 (B) TYPE: nucleic acid  
 (C) strandedness: double  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: human  
 (B) INDIVIDUAL ISOLATE: A549, ATCC CRL-185

(ix) FEATURE:

(A) NAME/KEY: mat peptide  
 (B) LOCATION: 1..522  
 (C) IDENTIFICATION METHOD: S

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 11:

TGC	GGA	CCG	GGC	AGG	GGG	TTC	GGG	AAA	AGG	AGG	CAC	CCC	AAA	AAG	CTG	48
Cys	Gly	Pro	Gly	Arg	Gly	Phe	Gly	Lys	Arg	Arg	His	Pro	Lys	Lys	Leu	
1				5				10						15		
ACC	CCT	TTA	GCC	TAC	AAG	CAG	TTT	ATC	CCC	AAT	GTG	GCC	GAA	AAG	ACC	96
Thr	Pro	Leu	Ala	Tyr	Lys	Gln	Phe	Ile	Pro	Asn	Val	Ala	Glu	Lys	Thr	
			20					25					30			
CTA	GGC	GCC	AGC	GGA	AGG	TAT	GAA	GGG	AAG	ATC	TCC	AGA	AAC	TCC	GAG	144
Leu	Gly	Ala	Ser	Gly	Arg	Tyr	Glu	Gly	Lys	Ile	Ser	Arg	Asn	Ser	Glu	

35					40					45										
CGA	TTT	AAG	GAA	CTC	ACC	CCC	AAT	TAC	AAC	CCC	GAC	ATC	ATA	TTT	AAG	192				
Arg	Phe	Lys	Glu	Leu	Thr	Pro	Asn	Tyr	Asn	Pro	Asp	Ile	Ile	Phe	Lys					
	50					55					60									
GAT	GAA	GAA	AAC	ACC	GGA	GCG	GAC	AGG	CTG	ATG	ACT	CAG	AGG	TGT	AAG	240				
Asp	Glu	Glu	Asn	Thr	Gly	Ala	Asp	Arg	Leu	Met	Thr	Gln	Arg	Cys	Lys					
	65				70					75					80					
GAC	AAG	TTG	AAC	GCT	TTG	GCC	ATC	TCG	GTG	ATG	AAC	CAG	TGG	CCA	GGA	288				
Asp	Lys	Leu	Asn	Ala	Leu	Ala	Ile	Ser	Val	Met	Asn	Gln	Trp	Pro	Gly					
				85					90					95						
GTG	AAA	CTG	CGG	GTG	ACC	GAG	GGC	TGG	GAC	GAA	GAT	GGC	CAC	CAC	TCA	336				
Val	Lys	Leu	Arg	Val	Thr	Glu	Gly	Trp	Asp	Glu	Asp	Gly	His	His	Ser					
			100					105					110							
GAG	GAG	TCT	CTG	CAC	TAC	GAG	GGC	CGC	GCA	GTG	GAC	ATC	ACC	ACG	TCT	384				
Glu	Glu	Ser	Leu	His	Tyr	Glu	Gly	Arg	Ala	Val	Asp	Ile	Thr	Thr	Ser					
		115					120					125								
GAC	CGC	GAC	CGC	AGC	AAG	TAC	GGC	ATG	CTG	GCC	CGC	CTG	GCG	GTG	GAG	432				
Asp	Arg	Asp	Arg	Ser	Lys	Tyr	Gly	Met	Leu	Ala	Arg	Leu	Ala	Val	Glu					
	130					135					140									
GCC	GGC	TTC	GAC	TGG	GTG	TAC	TAC	GAG	TCC	AAG	GCA	CAT	ATC	CAC	TGC	480				
Ala	Gly	Phe	Asp	Trp	Val	Tyr	Tyr	Glu	Ser	Lys	Ala	His	Ile	His	Cys					
	145				150					155					160					
TCG	GTG	AAA	GCA	GAG	AAC	TCG	GTG	GCG	GCC	AAA	TCG	GGA	GGC			522				
Ser	Val	Lys	Ala	Glu	Asn	Ser	Val	Ala	Ala	Lys	Ser	Gly	Gly							
				165				170					174							

(2) INFORMATION FOR SEQ ID NO: 12:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 20 base pairs  
(B) TYPE: nucleic acid  
(C) strandedness: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:

GCCAGGGTGT GAGCAACAGT

20

(2) INFORMATION FOR SEQ ID NO: 13:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 20 base pairs  
(B) TYPE: nucleic acid  
(C) strandedness: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:

TGTGCTGCTT GGCACCTCTTG

20

(2) INFORMATION FOR SEQ ID NO: 14:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 20 base pairs  
    (B) TYPE: nucleic acid  
    (C) strandedness: single  
    (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:

CCGTGGCATT TCCCGGAAAG

20

(2) INFORMATION FOR SEQ ID NO: 15:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 18 base pairs  
    (B) TYPE: nucleic acid  
    (C) strandedness: single  
    (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 15:

GSTATCCATGG CTCTCCTG

18

(2) INFORMATION FOR SEQ ID NO: 16:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 26 base pairs  
    (B) TYPE: nucleic acid  
    (C) strandedness: single  
    (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 16:

GCCTCGAGGT ATCCATGGCT CTCCTG

26

(2) INFORMATION FOR SEQ ID NO: 17:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 28 base pairs  
    (B) TYPE: nucleic acid  
    (C) strandedness: single  
    (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 17:

GCGCGGCCGC TCAGCCGCC GCGCGGAC

28

(2) INFORMATION FOR SEQ ID NO: 18:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 20 base pairs  
    (B) TYPE: nucleic acid  
    (C) strandedness: single  
    (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 18:

CGTGTCGGTC AAAGCTGATA

20

(2) INFORMATION FOR SEQ ID NO: 19:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 20 base pairs
  - (B) TYPE: nucleic acid
  - (C) strandedness: single
  - (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 19:

ATGCATTCCA GTCGGCTGGA

20

(2) INFORMATION FOR SEQ ID NO: 20:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 56 base pairs
  - (B) TYPE: nucleic acid
  - (C) strandedness: single
  - (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 20:

AAGGATCCGT CGACAAGCTT AATACGACGA ATTCTGGAGT TTTTTTTTTT TTTTTT

56

(2) INFORMATION FOR SEQ ID NO: 21:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 20 base pairs
  - (B) TYPE: nucleic acid
  - (C) strandedness: single
  - (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 21:

GGCTTCGACT GGGTCTACTA

20

(2) INFORMATION FOR SEQ ID NO: 22:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 17 base pairs
  - (B) TYPE: nucleic acid
  - (C) strandedness: single
  - (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:

AAGGATCCGT CGACAAG

17

(2) INFORMATION FOR SEQ ID NO: 23:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 17 base pairs
  - (B) TYPE: nucleic acid
  - (C) strandedness: single
  - (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 23:

ATGCGCTTCG GCCAGCG

17

(2) INFORMATION FOR SEQ ID NO: 24:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 17 base pairs
- (B) TYPE: nucleic acid
- (C) strandedness: single
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 24:

GACAAGCTTA ATACGAC

17

(2) INFORMATION FOR SEQ ID NO: 25:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 17 base pairs
- (B) TYPE: nucleic acid
- (C) strandedness: single
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 25:

GTTCGCGCCG CTCACCG

17

(2) INFORMATION FOR SEQ ID NO: 26:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 18 base pairs
- (B) TYPE: nucleic acid
- (C) strandedness: single
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 26:

TACGACGAAT TCTGGAGT

18

(2) INFORMATION FOR SEQ ID NO: 27:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 34 base pairs
- (B) TYPE: nucleic acid
- (C) strandedness: single
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 27:

CCCGGGAATT CATTGCGGGC CGGGCCGGGG GCCG

34

(2) INFORMATION FOR SEQ ID NO: 28:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 34 base pairs
- (B) TYPE: nucleic acid

(C) strandedness: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28:

ACGATGAATT CTCAGCCGCC CGCCCGGACC GCCA

34

(2) INFORMATION FOR SEQ ID NO: 29:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 6 amino acids  
(B) TYPE: amino acid  
(C) strandedness: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 29:

Gly Ser Pro Gly Ile His  
1 5

(2) INFORMATION FOR SEQ ID NO: 30:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 33 base pairs  
(B) TYPE: nucleic acid  
(C) strandedness: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 30:

CCCGGGAATT CATTGCGGAC CGGGCAGGGG GTT

33

(2) INFORMATION FOR SEQ ID NO: 31:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 32 base pairs  
(B) TYPE: nucleic acid  
(C) strandedness: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 31:

ACGATGAATT CTCAGCCTCC CGATTTGGCC GC

32



**WE CLAIM:**

1. A Desert hedgehog protein of human origin.
2. The hedgehog protein of claim 1, which contains a part or the whole of the amino acid sequence of SEQ ID NO:1.
3. The hedgehog protein of claim 1, which contains a part or the whole of the amino acid sequence of SEQ ID NO:2.
4. The hedgehog protein of claim 1, which contains a part or the whole of the amino acid sequence of SEQ ID NO:3.
5. The hedgehog protein of claim 1, which originates from a human cell.
6. The hedgehog protein of claim 1, which originates from established human cell line ARH-77, ATCC CRL-1621.
7. A DNA which encodes the hedgehog protein of claim 1.
8. The DNA of claim 7, which contains a part or the whole of either the nucleotide sequence of SEQ ID NO:4 or its complementary nucleotide sequence.
9. The DNA of claim 7, which contains a part or the whole of either the nucleotide sequence of SEQ ID NO:5 or its complementary nucleotide sequence.
10. The DNA of claim 7, which contains a part or the whole of either the nucleotide sequence of SEQ ID NO:6 or its complementary nucleotide sequence.
11. The DNA of claim 7, wherein, based on the degeneracy of genetic codes, one or more nucleotides are replaced with different nucleotides while conserving the encoding amino acid sequence.
12. The DNA of claim 7, which is inserted into an

autonomously replicable vector.

13. The DNA of claim 7, which is introduced into an appropriate host.

14. A monoclonal antibody which recognizes the hedgehog protein of claim 1.

15. The monoclonal antibody of claim 14, which additionally recognizes a Sonic hedgehog protein of human origin.

16. A hybridoma capable of producing a monoclonal antibody which recognizes the hedgehog protein of claim 1.

17. A process for producing a hedgehog protein which comprises the steps of allowing to express a DNA that encodes the hedgehog protein of claim 1 and collecting the generated hedgehog protein.

18. The process of claim 17, wherein the DNA is expressed through culturing of a transformant introduced with a DNA that encodes the hedgehog protein of claim 1.

19. The process of claim 17, wherein the generated hedgehog protein is collected by salting out, dialysis, filtration, concentration, fractional precipitation, ion-exchange chromatography, gel filtration chromatography, adsorption chromatography, isoelectric focusing chromatography, hydrophobic chromatography, reversed phase chromatography, affinity chromatography, gel electrophoresis and/or isoelectric focusing gel electrophoresis.

20. The process of claim 17, wherein the generated hedgehog protein is collected through immunoaffinity chromatography using a monoclonal antibody that recognizes a Desert hedgehog protein of human origin.

21. A method for detecting a hedgehog protein which

comprises the steps of bringing a monoclonal antibody that recognizes the hedgehog protein of claim 1 into contact with a sample and detecting the hedgehog protein based on an immunoreaction.

22. The method of claim 21, wherein the monoclonal antibody is labelled with a radioactive substance, enzyme and/or fluorescent substance.

Abstract of the Disclosure

Disclosed are a novel hedgehog protein, i.e., a Desert hedgehog protein of human origin including mature and precursor forms, a DNA encoding the protein, a monoclonal antibody recognizing the protein, a process for producing the protein, and a method for detecting the protein. The hedgehog protein is useful in establishment of hybridomas which produce antibodies recognizing the protein, and the monoclonal antibody is useful in detection and purification of the protein. The hedgehog protein, DNA, and monoclonal antibody of this invention have efficacy in elucidation of hereditary morphological abnormalities in humans to establish their treatments and diagnoses.

05067-0309

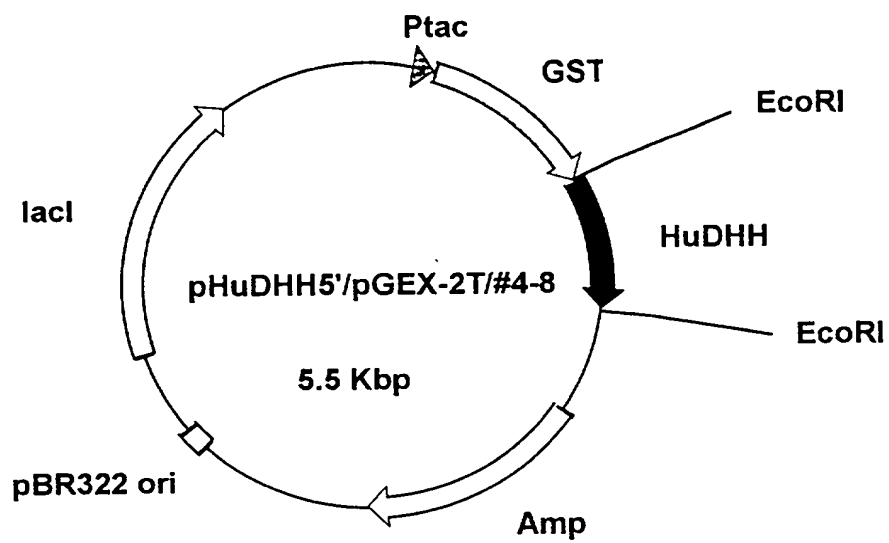
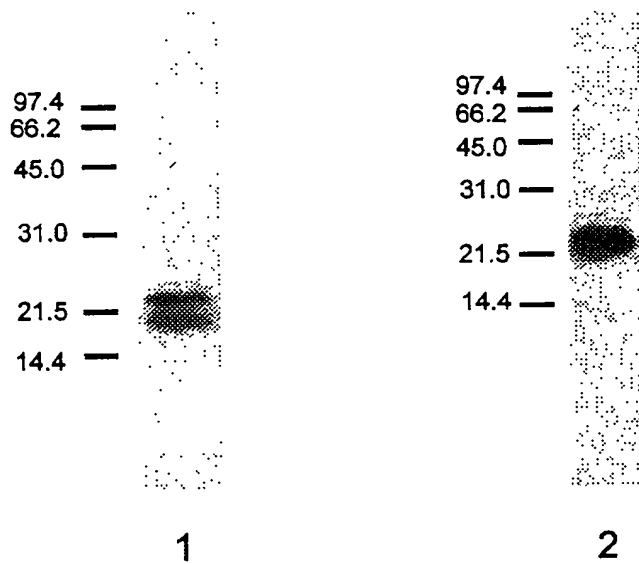


FIG.1



**FIG.2**

Note: On Lane 1, human Desert hedgehog protein was electrophoresed. On Lane 2, human Sonic hedgehog protein was electrophoresed. Numbers on left side of each lane mean the molecular weights of molecular weight markers in a unit of kilodaltons and indicate their positions after electrophoresis.

962140-8220060

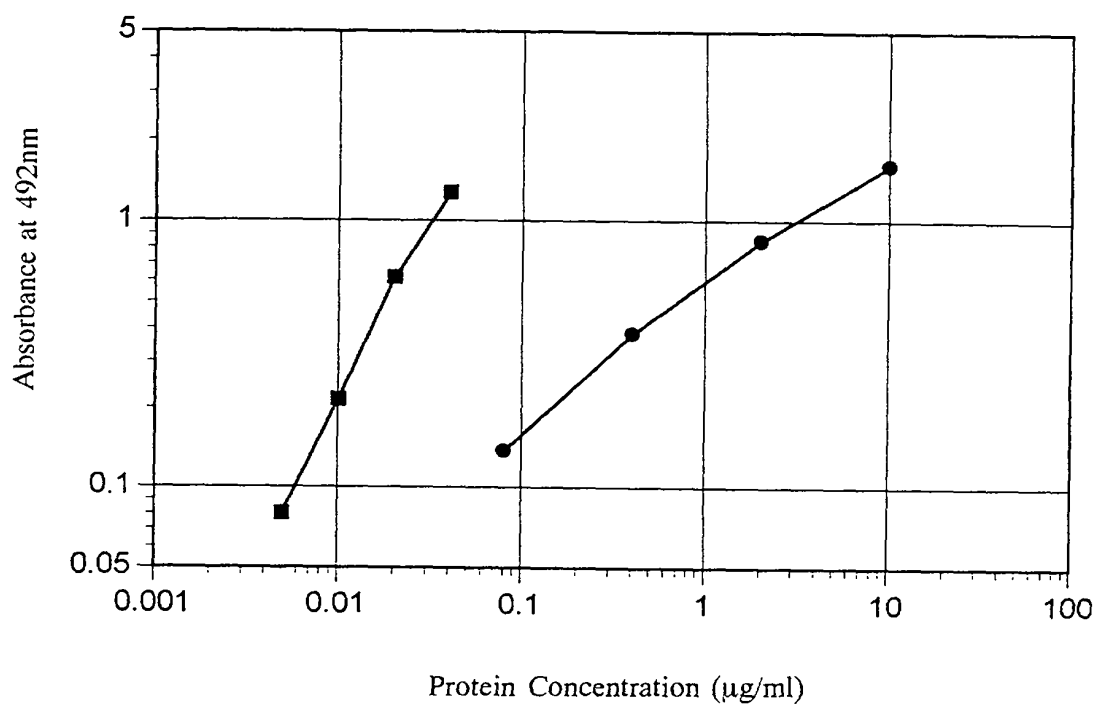


FIG.3

Note: Closed circles represent the results of detecting human Desert hedgehog protein, and closed squares represent the results of detecting human Sonic hedgehog protein.

The undersigned hereby authorizes the U.S. Attorneys or Agents named herein to accept and follow instructions from SUMA PATENT OFFICE as to any action to be taken in the U.S. Patent and Trademark Office regarding this application without direct communication between the U.S. Attorney or Agent and the undersigned. In the event of a change of the persons from whom instructions may be taken, the U.S. Attorneys or Agents named herein will be so notified by the undersigned.



Page 2 of 2      **Atty.Docket:**  
 Title: HEDGEHOG PROTEIN  
 U.S. Application filed \_\_\_\_\_, Serial No. \_\_\_\_\_  
 PCT Application filed \_\_\_\_\_, Serial No. \_\_\_\_\_

I hereby further declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. § 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

FULL NAME OF FIRST INVENTOR Toshio ARIYASU		INVENTOR'S SIGNATURE <i>Toshio Ariyasu</i>	DATE <i>April 15, 1998</i>
RESIDENCE Okayama, Japan		CITIZENSHIP Japanese	
POST OFFICE ADDRESS 427-2, Ohfuku, Okayama-shi, Okayama, Japan			
FULL NAME OF SECOND JOINT INVENTOR Shuji NAKAMURA		INVENTOR'S SIGNATURE <i>Shuji Nakamura</i>	DATE <i>April 15, 1998</i>
RESIDENCE Okayama, Japan		CITIZENSHIP Japanese	
POST OFFICE ADDRESS 525-3, Kuwano, Okayama-shi, Okayama, Japan			
FULL NAME OF THIRD JOINT INVENTOR Kunzo ORITA		INVENTOR'S SIGNATURE <i>Kunzo Orita</i>	DATE <i>April 15, 1998</i>
RESIDENCE Okayama, Japan		CITIZENSHIP Japanese	
POST OFFICE ADDRESS 2-34, 1-chome, Nishizaka, Tsushima, Okayama-shi, Okayama, Japan			
FULL NAME OF FOURTH JOINT INVENTOR		INVENTOR'S SIGNATURE	DATE
RESIDENCE		CITIZENSHIP	
POST OFFICE ADDRESS			
FULL NAME OF FIFTH JOINT INVENTOR		INVENTOR'S SIGNATURE	DATE
RESIDENCE		CITIZENSHIP	
POST OFFICE ADDRESS			
FULL NAME OF SIXTH JOINT INVENTOR		INVENTOR'S SIGNATURE	DATE
RESIDENCE		CITIZENSHIP	
POST OFFICE ADDRESS			
FULL NAME OF SEVENTH JOINT INVENTOR		INVENTOR'S SIGNATURE	DATE
RESIDENCE		CITIZENSHIP	
POST OFFICE ADDRESS			

ALL INVENTORS MUST REVIEW APPLICATION AND DECLARATION BEFORE SIGNING. ALL ALTERATIONS MUST BE INITIALED AND DATED BY ALL INVENTORS PRIOR TO EXECUTION. NO ALTERATIONS CAN BE MADE AFTER THE DECLARATION IS SIGNED. ALL PAGES OF DECLARATION MUST BE SEEN BY ALL INVENTORS.

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